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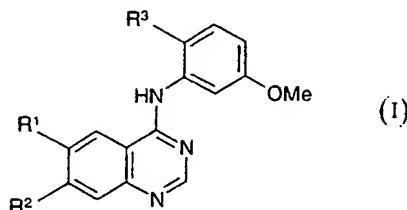
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(54) Title: QUINAZOLINE DERIVATIVES



(57) Abstract: The invention concerns quinazoline derivatives of Formula (I), wherein each of R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> have any of the meanings defined in the description; processes for their preparation, pharmaceutical compositions containing them and their use in the manufacture of a medicament for use as an anti-invasive agent in the containment and/or treatment of solid tumour disease.



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QUINAZOLINE DERIVATIVES

The invention concerns certain novel quinazoline derivatives, or pharmaceutically-acceptable salts thereof, which possess anti-tumour activity and are  
5 accordingly useful in methods of treatment of the human or animal body. The invention also concerns processes for the manufacture of said quinazoline derivatives, to pharmaceutical compositions containing them and to their use in therapeutic methods, for example in the manufacture of medicaments for use in the prevention or treatment of solid tumour disease in a warm-blooded animal such as man.

10 Many of the current treatment regimes for cell proliferation diseases such as psoriasis and cancer utilise compounds which inhibit DNA synthesis. Such compounds are toxic to cells generally but their toxic effect on rapidly dividing cells such as tumour cells can be beneficial. Alternative approaches to anti-tumour agents which act by mechanisms other than the inhibition of DNA synthesis have the potential to display enhanced selectivity of action.

15 In recent years it has been discovered that a cell may become cancerous by virtue of the transformation of a portion of its DNA into an oncogene i.e. a gene which, on activation, leads to the formation of malignant tumour cells (Bradshaw, Mutagenesis, 1986, 1, 91). Several such oncogenes give rise to the production of peptides which are receptors for growth factors. Activation of the growth factor receptor complex subsequently leads to an increase in  
20 cell proliferation. It is known, for example, that several oncogenes encode tyrosine kinase enzymes and that certain growth factor receptors are also tyrosine kinase enzymes (Yarden *et al.*, Ann. Rev. Biochem., 1988, 57, 443; Larsen *et al.*, Ann. Reports in Med. Chem., 1989, Chpt. 13). The first group of tyrosine kinases to be identified arose from such viral oncogenes, for example pp60<sup>v-Src</sup> tyrosine kinase (otherwise known as v-Src), and the  
25 corresponding tyrosine kinases in normal cells, for example pp60<sup>c-Src</sup> tyrosine kinase (otherwise known as c-Src).

Receptor tyrosine kinases are important in the transmission of biochemical signals which initiate cell replication. They are large enzymes which span the cell membrane and possess an extracellular binding domain for growth factors such as epidermal growth factor  
30 (EGF) and an intracellular portion which functions as a kinase to phosphorylate tyrosine amino acids in proteins and hence to influence cell proliferation. Various classes of receptor tyrosine kinases are known (Wilks, Advances in Cancer Research, 1993, 60, 43-73) based on families of growth factors which bind to different receptor tyrosine kinases. The classification

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includes Class I receptor tyrosine kinases comprising the EGF family of receptor tyrosine kinases such as the EGF, TGF $\alpha$ , Neu and erbB receptors, Class II receptor tyrosine kinases comprising the insulin family of receptor tyrosine kinases such as the insulin and IGF1 receptors and insulin-related receptor (IRR) and Class III receptor tyrosine kinases comprising the platelet-derived growth factor (PDGF) family of receptor tyrosine kinases such as the PDGF $\alpha$ , PDGF $\beta$  and colony-stimulating factor 1 (CSF1) receptors.

It is also known that certain tyrosine kinases belong to the class of non-receptor tyrosine kinases which are located intracellularly and are involved in the transmission of biochemical signals such as those that influence tumour cell motility, dissemination and invasiveness and subsequently metastatic tumour growth (Ullrich *et al.*, Cell, 1990, 61, 203-212, Bolen *et al.*, FASEB J., 1992, 6, 3403-3409, Brickell *et al.*, Critical Reviews in Oncogenesis, 1992, 3, 401-406, Bohlen *et al.*, Oncogene, 1993, 8, 2025-2031, Courtneidge *et al.*, Semin. Cancer Biol., 1994, 5, 239-246, Lauffenburger *et al.*, Cell, 1996, 84, 359-369, Hanks *et al.*, BioEssays, 1996, 19, 137-145, Parsons *et al.*, Current Opinion in Cell Biology, 1997, 9, 187-192, Brown *et al.*, Biochimica et Biophysica Acta, 1996, 1287, 121-149 and Schlaepfer *et al.*, Progress in Biophysics and Molecular Biology, 1999, 71, 435-478). Various classes of non-receptor tyrosine kinases are known including the Src family such as the Src, Lyn, Fyn and Yes tyrosine kinases, the Abl family such as Abl and Arg and the Jak family such as Jak 1 and Tyk 2.

It is known that the Src family of non-receptor tyrosine kinases are highly regulated in normal cells and in the absence of extracellular stimuli are maintained in an inactive conformation. However, some Src family members, for example c-Src tyrosine kinase, is frequently significantly activated (when compared to normal cell levels) in common human cancers such as gastrointestinal cancer, for example colon, rectal and stomach cancer (Cartwright *et al.*, Proc. Natl. Acad. Sci. USA, 1990, 87, 558-562 and Mao *et al.*, Oncogene, 1997, 15, 3083-3090), and breast cancer (Muthuswamy *et al.*, Oncogene, 1995, 11, 1801-1810). The Src family of non-receptor tyrosine kinases has also been located in other common human cancers such as non-small cell lung cancers (NSCLCs) including adenocarcinomas and squamous cell cancer of the lung (Mazurenko *et al.*, European Journal of Cancer, 1992, 28, 372-7), bladder cancer (Fanning *et al.*, Cancer Research, 1992, 52, 1457-62), oesophageal cancer (Jankowski *et al.*, Gut, 1992, 33, 1033-8), cancer of the prostate, ovarian cancer (Wiener *et al.*, Clin. Cancer Research, 1999, 5, 2164-70) and pancreatic cancer

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(Lutz *et al.*, Biochem. and Biophys. Res. Comm., 1998, 243, 503-8). As further human tumour tissues are tested for the Src family of non-receptor tyrosine kinases it is expected that its widespread prevalence will be established.

It is further known that the predominant role of c-Src non-receptor tyrosine kinase is to  
5 regulate the assembly of focal adhesion complexes through interaction with a number of  
cytoplasmic proteins including, for example, focal adhesion kinase and paxillin. In addition  
c-Src is coupled to signalling pathways that regulate the actin cytoskeleton which facilitates  
cell motility. Likewise, important roles are played by the c-Src, c-Yes and c-Fyn non-receptor  
tyrosine kinases in integrin mediated signalling and in disrupting cadherin-dependent cell-cell  
10 junctions (Owens *et al.*, Molecular Biology of the Cell, 2000, 11, 51-64 and Klinghoffer *et al.*,  
EMBO Journal, 1999, 18, 2459-2471). Cellular motility is necessarily required for a localised  
tumour to progress through the stages of dissemination into the blood stream, invasion of  
other tissues and initiation of metastatic tumour growth. For example, colon tumour  
progression from localised to disseminated, invasive metastatic disease has been correlated  
15 with c-Src non-receptor tyrosine kinase activity (Brunton *et al.*, Oncogene, 1997, 14, 283-293,  
Fincham *et al.*, EMBO J, 1998, 17, 81-92 and Verbeek *et al.*, Exp. Cell Research, 1999, 248,  
531-537).

Accordingly it has been recognised that an inhibitor of such non-receptor tyrosine  
kinases should be of value as a selective inhibitor of the motility of tumour cells and as a  
20 selective inhibitor of the dissemination and invasiveness of mammalian cancer cells leading to  
inhibition of metastatic tumour growth. In particular an inhibitor of such non-receptor  
tyrosine kinases should be of value as an anti-invasive agent for use in the containment and/or  
treatment of solid tumour disease.

We have now found that surprisingly certain quinazoline derivatives possess potent  
25 anti-tumour activity. Without wishing to imply that the compounds disclosed in the present  
invention possess pharmacological activity only by virtue of an effect on a single biological  
process, it is believed that the compounds provide an anti-tumour effect by way of inhibition  
of one or more of the non-receptor tyrosine-specific protein kinases that are involved in the  
signal transduction steps which lead to the invasiveness and migratory ability of metastasising  
30 tumour cells. In particular, it is believed that the compounds of the present invention provide  
an anti-tumour effect by way of inhibition of the Src family of non-receptor tyrosine kinases,  
for example by inhibition of one or more of c-Src, c-Yes and c-Fyn.



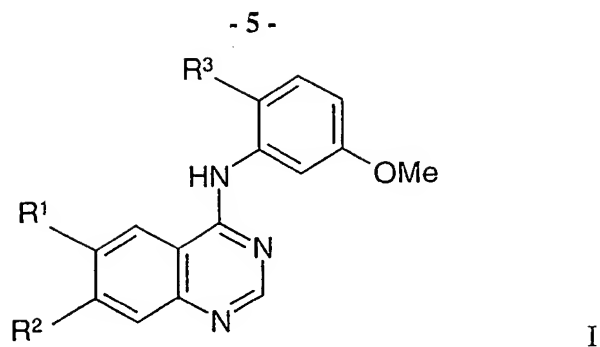
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It is also known that c-Src non-receptor tyrosine kinase enzyme is involved in the control of osteoclast-driven bone resorption (Soriano *et al.*, Cell, 1991, 64, 693-702; Boyce *et al.*, J. Clin. Invest., 1992, 90, 1622-1627; Yoneda *et al.*, J. Clin. Invest., 1993, 91, 2791-2795 and Missbach *et al.*, Bone, 1999, 24, 437-49). An inhibitor of c-Src non-receptor tyrosine  
5 kinase is therefore of value in the prevention and treatment of bone diseases such as osteoporosis, Paget's disease, metastatic disease in bone and tumour-induced hypercalcaemia.

The compounds of the present invention are also useful in inhibiting the uncontrolled cellular proliferation which arises from various non-malignant diseases such as inflammatory  
10 diseases (for example rheumatoid arthritis and inflammatory bowel disease), fibrotic diseases (for example hepatic cirrhosis and lung fibrosis), glomerulonephritis, multiple sclerosis, psoriasis, hypersensitivity reactions of the skin, blood vessel diseases (for example atherosclerosis and restenosis), allergic asthma, insulin-dependent diabetes, diabetic retinopathy and diabetic nephropathy.

15 Generally the compounds of the present invention possess potent inhibitory activity against the Src family of non-receptor tyrosine kinases, for example by inhibition of c-Src and/or c-Yes, whilst possessing less potent inhibitory activity against other tyrosine kinase enzymes such as the receptor tyrosine kinases, for example EGF receptor tyrosine kinase and/or VEGF receptor tyrosine kinase. Furthermore, certain compounds of the present  
20 invention possess substantially better potency against the Src family of non-receptor tyrosine kinases, for example c-Src and/or c-Yes, than against VEGF receptor tyrosine kinase. Such compounds possess sufficient potency against the Src family of non-receptor tyrosine kinases, for example c-Src and/or c-Yes, that they may be used in an amount sufficient to inhibit, for example, c-Src and/or c-Yes whilst demonstrating little activity against VEGF receptor  
25 tyrosine kinase.

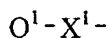
According to one aspect of the invention there is provided a quinazoline derivative of the Formula I



wherein :-

$R^3$  is chloro, bromo or iodo;

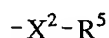
5  $R^1$  is hydrogen or (1-6C)alkoxy and  $R^2$  is a group of the formula :



wherein  $X^1$  is selected from O, S, SO, SO<sub>2</sub>, N(R<sup>4</sup>), CO, CH(OR<sup>4</sup>), CON(R<sup>4</sup>), N(R<sup>4</sup>)CO, SO<sub>2</sub>N(R<sup>4</sup>), N(R<sup>4</sup>)SO<sub>2</sub>, OC(R<sup>4</sup>)<sub>2</sub>, SC(R<sup>4</sup>)<sub>2</sub> and N(R<sup>4</sup>)C(R<sup>4</sup>)<sub>2</sub>, wherein R<sup>4</sup> is hydrogen or (1-6C)alkyl, and Q<sup>1</sup> is heteroaryl, heteroaryl-(1-6C)alkyl, heteroaryloxy-(1-6C)alkyl,

10 heterocyclyl, heterocyclyl-(1-6C)alkyl or heterocyclyloxy-(1-6C)alkyl, provided that, when X<sup>1</sup> is O, any compound wherein Q<sup>1</sup> is a piperidino-(2-4C)alkyl, morpholino-(2-4C)alkyl or piperazin-1-yl-(2-4C)alkyl group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is excluded, and provided that, when X<sup>1</sup> is O, any compound wherein Q<sup>1</sup> is a 4-(1-4C)alkylpiperazin-1-yl-  
 15 (2-4C)alkyl group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is also excluded,

or  $R^2$  is a group of the formula :



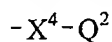
wherein X<sup>2</sup> is selected from O and N(R<sup>6</sup>), wherein R<sup>6</sup> is hydrogen or (1-6C)alkyl, and R<sup>5</sup> is  
 20 hydroxy-(3-6C)alkyl, (1-6C)alkoxy-(3-6C)alkyl, amino-(3-6C)alkyl, (1-6C)alkylamino-(3-6C)alkyl, di-[(1-6C)alkyl]amino-(3-6C)alkyl, (2-6C)alkanoylamino-(3-6C)alkyl or (1-6C)alkoxycarbonylamino-(3-6C)alkyl, provided that the central CH<sub>2</sub> group within any (CH<sub>2</sub>)<sub>3</sub> group within the (3-6C)alkylene group in a R<sup>5</sup> group bears a substituent selected from hydroxy, amino, (1-6C)alkoxy, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (2-6C)alkanoyloxy,  
 25 (2-6C)alkanoylamino and N-(1-6C)alkyl-(2-6C)alkanoylamino,

and wherein adjacent carbon atoms in any (2-6C)alkylene chain within a R<sup>2</sup> substituent are optionally separated by the insertion into the chain of a group selected from O, S, SO, SO<sub>2</sub>,

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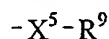
$N(R^7)$ ,  $CO$ ,  $CH(OR^7)$ ,  $CON(R^7)$ ,  $N(R^7)CO$ ,  $SO_2N(R^7)$ ,  $N(R^7)SO_2$ ,  $CH=CH$  and  $C\equiv C$  wherein  $R^7$  is hydrogen or (1-6C)alkyl, or, when the inserted group is  $N(R^7)$ ,  $R^7$  may also be (2-6C)alkanoyl,

and wherein any  $CH_2$  or  $CH_3$  group within a  $R^2$  substituent optionally bears on each  
 5 said  $CH_2$  or  $CH_3$  group one or more halogeno or (1-6C)alkyl substituents or a substituent selected from hydroxy, cyano, amino, carboxy, carbamoyl, (2-8C)alkenyl, (2-8C)alkynyl, (1-6C)alkoxy, (1-6C)alkylthio, (1-6C)alkylsulphinyl, (1-6C)alkylsulphonyl, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (1-6C)alkoxycarbonyl, N-(1-6C)alkylcarbamoyl, N,N-di-[(1-6C)alkyl]carbamoyl, (2-6C)alkanoyl, (2-6C)alkanoyloxy, (2-6C)alkanoylamino,  
 10 N-(1-6C)alkyl-(2-6C)alkanoylamino, N-(1-6C)alkylsulphamoyl, N,N-di-[(1-6C)alkyl]sulphamoyl, (1-6C)alkanesulphonylamino and N-(1-6C)alkyl-(1-6C)alkanesulphonylamino, or from a group of the formula :



wherein  $X^4$  is a direct bond or is selected from O, S, SO,  $SO_2$ ,  $N(R^8)$ , CO,  $CH(OR^8)$ ,  
 15  $CON(R^8)$ ,  $N(R^8)CO$ ,  $SO_2N(R^8)$ ,  $N(R^8)SO_2$ ,  $C(R^8)_2O$ ,  $C(R^8)_2S$  and  $N(R^8)C(R^8)_2$ , wherein  $R^8$  is hydrogen or (1-6C)alkyl, and  $Q^2$  is aryl, aryl-(1-6C)alkyl, (3-7C)cycloalkyl, (3-7C)cycloalkyl-(1-6C)alkyl, (3-7C)cycloalkenyl, (3-7C)cycloalkenyl-(1-6C)alkyl, heteroaryl, heteroaryl-(1-6C)alkyl, heterocyclyl or heterocyclyl-(1-6C)alkyl,

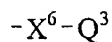
and wherein any aryl, heteroaryl or heterocyclyl group within a substituent on  $R^2$   
 20 optionally bears 1, 2 or 3 substituents, which may be the same or different, selected from halogeno, trifluoromethyl, cyano, nitro, hydroxy, amino, carboxy, carbamoyl, (1-6C)alkyl, (2-8C)alkenyl, (2-8C)alkynyl, (1-6C)alkoxy, (2-6C)alkenyloxy, (2-6C)alkynyloxy, (1-6C)alkylthio, (1-6C)alkylsulphinyl, (1-6C)alkylsulphonyl, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (1-6C)alkoxycarbonyl, N-(1-6C)alkylcarbamoyl,  
 25 N,N-di-[(1-6C)alkyl]carbamoyl, (2-6C)alkanoyl, (2-6C)alkanoyloxy, (2-6C)alkanoylamino, N-(1-6C)alkyl-(2-6C)alkanoylamino, N-(1-6C)alkylsulphamoyl, N,N-di-[(1-6C)alkyl]sulphamoyl, (1-6C)alkanesulphonylamino and N-(1-6C)alkyl-(1-6C)alkanesulphonylamino, or from a group of the formula :



30 wherein  $X^5$  is a direct bond or is selected from O and  $N(R^{10})$ , wherein  $R^{10}$  is hydrogen or (1-6C)alkyl, and  $R^9$  is halogeno-(1-6C)alkyl, hydroxy-(1-6C)alkyl, (1-6C)alkoxy-(1-6C)alkyl, cyano-(1-6C)alkyl, amino-(1-6C)alkyl, (1-6C)alkylamino-(1-6C)alkyl, di-[(1-6C)alkyl]amino-

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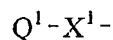
(1-6C)alkyl, (2-6C)alkanoylamino-(1-6C)alkyl or (1-6C)alkoxycarbonylamino-(1-6C)alkyl, or from a group of the formula :



wherein  $X^6$  is a direct bond or is selected from O, CO and  $N(R^{11})$ , wherein  $R^{11}$  is hydrogen or  
 5 (1-6C)alkyl, and  $Q^3$  is aryl, aryl-(1-6C)alkyl, heteroaryl, heteroaryl-(1-6C)alkyl, heterocyclyl or heterocyclyl-(1-6C)alkyl which optionally bears 1 or 2 substituents, which may be the same or different, selected from halogeno, (1-6C)alkyl, (2-8C)alkenyl, (2-8C)alkynyl and (1-6C)alkoxy,

and wherein any heterocyclyl group within a substituent on  $R^2$  optionally bears 1 or 2  
 10 oxo or thioxo substituents;

or wherein  $R^2$  is hydrogen or (1-6C)alkoxy and  $R^1$  is a group of the formula :



wherein  $X^1$  is selected from O, S, SO,  $SO_2$ ,  $N(R^4)$ , CO,  $CH(OR^4)$ ,  $CON(R^4)$ ,  $N(R^4)CO$ ,  
 15  $SO_2N(R^4)$ ,  $N(R^4)SO_2$ ,  $OC(R^4)_2$ ,  $SC(R^4)_2$  and  $N(R^4)C(R^4)_2$ , wherein  $R^4$  is hydrogen or (1-6C)alkyl, and  $Q^1$  is heterocyclyl, heterocyclyl-(1-6C)alkyl or heterocyclyoxy-(1-6C)alkyl, provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a piperidino-(2-4C)alkyl, morpholino-(2-4C)alkyl or piperazin-1-yl-(2-4C)alkyl group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the (2-4C)alkylene  
 20 group is excluded, and provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a 4-(1-4C)alkylpiperazin-1-yl-(2-4C)alkyl group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is also excluded,

or  $R^1$  is a group of the formula :



wherein  $X^2$  is selected from O and  $N(R^6)$ , wherein  $R^6$  is hydrogen or (1-6C)alkyl, and  $R^5$  is hydroxy-(3-6C)alkyl, (1-6C)alkoxy-(3-6C)alkyl, amino-(3-6C)alkyl, (1-6C)alkylamino-(3-6C)alkyl, di-[(1-6C)alkyl]amino-(3-6C)alkyl, (2-6C)alkanoylamino-(3-6C)alkyl or (1-6C)alkoxycarbonylamino-(3-6C)alkyl, provided that the central  $CH_2$  group within any  
 30  $(CH_2)_3$  group within the (3-6C)alkylene group in a  $R^5$  group bears a substituent selected from hydroxy, amino, (1-6C)alkoxy, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (2-6C)alkanoyloxy, (2-6C)alkanoylamino and  $\underline{N}$ -(1-6C)alkyl-(2-6C)alkanoylamino,

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and wherein adjacent carbon atoms in any (2-6C)alkylene chain within a  $R^1$  substituent are optionally separated by the insertion into the chain of a group selected from O, S, SO, SO<sub>2</sub>, N(R<sup>7</sup>), CO, CH(OR<sup>7</sup>), CON(R<sup>7</sup>), N(R<sup>7</sup>)CO, SO<sub>2</sub>N(R<sup>7</sup>), N(R<sup>7</sup>)SO<sub>2</sub>, CH=CH and C≡C wherein R<sup>7</sup> is hydrogen or (1-6C)alkyl, or, when the inserted group is N(R<sup>7</sup>), R<sup>7</sup> may also be

5 (2-6C)alkanoyl,

and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a  $R^1$  substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group one or more halogeno or (1-6C)alkyl substituents or a substituent selected from hydroxy, cyano, amino, carboxy, carbamoyl, (2-8C)alkenyl, (2-8C)alkynyl, (1-6C)alkoxy, (1-6C)alkylthio, (1-6C)alkylsulphinyl, (1-6C)alkylsulphonyl, (1-6C)alkylamino, 10 di-[(1-6C)alkyl]amino, (1-6C)alkoxycarbonyl, N-(1-6C)alkylcarbamoyl, N,N-di-[(1-6C)alkyl]carbamoyl, (2-6C)alkanoyl, (2-6C)alkanoyloxy, (2-6C)alkanoylamino, N-(1-6C)alkyl-(2-6C)alkanoylamino, N-(1-6C)alkylsulphamoyl, N,N-di-[(1-6C)alkyl]sulphamoyl, (1-6C)alkanesulphonylamino and N-(1-6C)alkyl-(1-6C)alkanesulphonylamino, or from a group of the formula :

15  $-X^4-Q^2$

wherein  $X^4$  is a direct bond or is selected from O, S, SO, SO<sub>2</sub>, N(R<sup>8</sup>), CO, CH(OR<sup>8</sup>), CON(R<sup>8</sup>), N(R<sup>8</sup>)CO, SO<sub>2</sub>N(R<sup>8</sup>), N(R<sup>8</sup>)SO<sub>2</sub>, C(R<sup>8</sup>)<sub>2</sub>O, C(R<sup>8</sup>)<sub>2</sub>S and N(R<sup>8</sup>)C(R<sup>8</sup>)<sub>2</sub>, wherein R<sup>8</sup> is hydrogen or (1-6C)alkyl, and Q<sup>2</sup> is aryl, aryl-(1-6C)alkyl, (3-7C)cycloalkyl, (3-7C)cycloalkyl-(1-6C)alkyl, (3-7C)cycloalkenyl, (3-7C)cycloalkenyl-(1-6C)alkyl, heteroaryl, heteroaryl- 20 (1-6C)alkyl, heterocyclyl or heterocyclyl-(1-6C)alkyl,

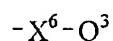
and wherein any aryl, heteroaryl or heterocyclyl group within a substituent on  $R^1$  optionally bears 1, 2 or 3 substituents, which may be the same or different, selected from halogeno, trifluoromethyl, cyano, nitro, hydroxy, amino, carboxy, carbamoyl, (1-6C)alkyl, (2-8C)alkenyl, (2-8C)alkynyl, (1-6C)alkoxy, (2-6C)alkenyloxy, (2-6C)alkynyloxy, 25 (1-6C)alkylthio, (1-6C)alkylsulphinyl, (1-6C)alkylsulphonyl, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (1-6C)alkoxycarbonyl, N-(1-6C)alkylcarbamoyl, N,N-di-[(1-6C)alkyl]carbamoyl, (2-6C)alkanoyl, (2-6C)alkanoyloxy, (2-6C)alkanoylamino, N-(1-6C)alkyl-(2-6C)alkanoylamino, N-(1-6C)alkylsulphamoyl, N,N-di-[(1-6C)alkyl]sulphamoyl, (1-6C)alkanesulphonylamino and N-(1-6C)alkyl- 30 (1-6C)alkanesulphonylamino, or from a group of the formula :

$-X^5-R^9$

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wherein  $X^5$  is a direct bond or is selected from O and  $N(R^{10})$ , wherein  $R^{10}$  is hydrogen or (1-6C)alkyl, and  $R^9$  is halogeno-(1-6C)alkyl, hydroxy-(1-6C)alkyl, (1-6C)alkoxy-(1-6C)alkyl, cyano-(1-6C)alkyl, amino-(1-6C)alkyl, (1-6C)alkylamino-(1-6C)alkyl, di-[(1-6C)alkyl]amino-(1-6C)alkyl, (2-6C)alkanoylamino-(1-6C)alkyl or (1-6C)alkoxycarbonylamino-(1-6C)alkyl,

5 or from a group of the formula :



wherein  $X^6$  is a direct bond or is selected from O, CO and  $N(R^{11})$ , wherein  $R^{11}$  is hydrogen or (1-6C)alkyl, and  $Q^3$  is aryl, aryl-(1-6C)alkyl, heteroaryl, heteroaryl-(1-6C)alkyl, heterocyclyl or heterocyclyl-(1-6C)alkyl which optionally bears 1 or 2 substituents, which may be the same  
10 or different, selected from halogeno, (1-6C)alkyl, (2-8C)alkenyl, (2-8C)alkynyl and (1-6C)alkoxy,

and wherein any heterocyclyl group within a substituent on  $R^1$  optionally bears 1 or 2 oxo or thioxo substituents;  
or a pharmaceutically-acceptable salt thereof.

15

In this specification the generic term "alkyl" includes both straight-chain and branched-chain alkyl groups such as propyl, isopropyl and tert-butyl, and (3-7C)cycloalkyl groups such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl. However references to individual alkyl groups such as "propyl" are specific for the straight-chain  
20 version only, references to individual branched-chain alkyl groups such as "isopropyl" are specific for the branched-chain version only and references to individual cycloalkyl groups such as "cyclopentyl" are specific for that 5-membered ring only. An analogous convention applies to other generic terms, for example (1-6C)alkoxy includes methoxy, ethoxy, cyclopropyloxy and cyclopentyloxy, (1-6C)alkylamino includes methylamino, ethylamino,  
25 cyclobutylamino and cyclohexylamino, and di-[(1-6C)alkyl]amino includes dimethylamino, diethylamino, N-cyclobutyl-N-methylamino and N-cyclohexyl-N-ethylamino.

It is to be understood that, insofar as certain of the compounds of Formula I defined above may exist in optically active or racemic forms by virtue of one or more asymmetric carbon atoms, the invention includes in its definition any such optically active or racemic form  
30 which possesses the above-mentioned activity. The synthesis of optically active forms may be carried out by standard techniques of organic chemistry well known in the art, for example by synthesis from optically active starting materials or by resolution of a racemic form.

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Similarly, the above-mentioned activity may be evaluated using the standard laboratory techniques referred to hereinafter.

Suitable values for the generic radicals referred to above include those set out below.

A suitable value for  $Q^2$  or  $Q^3$  when it is aryl or for the aryl group within a 'Q' group is, for example, phenyl or naphthyl, conveniently phenyl.

A suitable value for  $Q^2$  when it is (3-7C)cycloalkyl or for the (3-7C)cycloalkyl group within a 'Q' group is, for example, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl or bicyclo[2.2.1]heptyl and a suitable value for  $Q^2$  when it is (3-7C)cycloalkenyl or for the (3-7C)cycloalkenyl group within a 'Q' group is, for example, cyclobutenyl, cyclopentenyl, cyclohexenyl or cycloheptenyl.

A suitable value for any one of the 'Q' groups ( $Q^1$  to  $Q^3$ ) when it is heteroaryl or for the heteroaryl group within a 'Q' group is, for example, an aromatic 5- or 6-membered monocyclic ring or a 9- or 10-membered bicyclic ring with up to five ring heteroatoms selected from oxygen, nitrogen and sulphur, for example furyl, pyrrolyl, thienyl, oxazolyl, isoxazolyl, imidazolyl, pyrazolyl, thiazolyl, isothiazolyl, oxadiazolyl, thiadiazolyl, triazolyl, tetrazolyl, pyridyl, pyridazinyl, pyrimidinyl, pyrazinyl, 1,3,5-triazenyl, benzofuranyl, indolyl, benzothienyl, benzoxazolyl, benzimidazolyl, benzothiazolyl, indazolyl, benzofurazanyl, quinolyl, isoquinolyl, quinazolinyl, quinoxalinyl, cinnolinyl or naphthyridinyl.

A suitable value for any one of the 'Q' groups ( $Q^1$  to  $Q^3$ ) when it is heterocyclyl or for the heterocyclyl group within a 'Q' group is, for example, a non-aromatic saturated or partially saturated 3 to 10 membered monocyclic or bicyclic ring with up to five heteroatoms selected from oxygen, nitrogen and sulphur, for example oxiranyl, oxetanyl, tetrahydrofuranyl, tetrahydropyranyl, oxepanyl, azetidiny, pyrrolinyl, pyrrolidinyl, morpholinyl, tetrahydro-1,4-thiazinyl, 1,1-dioxotetrahydro-1,4-thiazinyl, piperidinyl, homopiperidinyl, piperazinyl, homopiperazinyl, dihydropyridinyl, tetrahydropyridinyl, dihydropyrimidinyl or tetrahydropyrimidinyl, conveniently pyrrolidinyl, morpholinyl, 1,1-dioxotetrahydro-1,4-thiazinyl, piperidinyl, homopiperidinyl or piperazinyl. A suitable value for such a group which bears 1 or 2 oxo or thioxo substituents is, for example, 2-oxopyrrolidinyl, 2-thioxopyrrolidinyl, 2-oxoimidazolidinyl, 2-thioxoimidazolidinyl, 2-oxopiperidinyl, 2,5-dioxopyrrolidinyl, 2,5-dioxoimidazolidinyl or 2,6-dioxopiperidinyl.

A suitable value for a 'Q' group when it is heteroaryl-(1-6C)alkyl is, for example, heteroarylmethyl, 2-heteroarylethyl and 3-heteroarylpropyl. The invention comprises corresponding suitable values for 'Q' groups when, for example, rather than a

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heteroaryl-(1-6C)alkyl group, a heteroaryloxy-(1-6C)alkyl, an aryl-(1-6C)alkyl, (3-7C)cycloalkyl-(1-6C)alkyl, (3-7C)cycloalkenyl-(1-6C)alkyl, heterocyclyl-(1-6C)alkyl or heterocyclyloxy-(1-6C)alkyl group is present.

- Suitable values for any of the 'R' groups ( $R^1$  to  $R^{11}$ ), or for various groups within an
- 5  $R^1$  or  $R^2$  substituent, or for various groups within  $Q^1$ ,  $Q^2$  or  $Q^3$  include :-
- |  |   |
|--|---|
| for halogeno                                   | fluoro, chloro, bromo and iodo;   |
| for (1-6C)alkyl:                               | methyl, ethyl, propyl, isopropyl and <u>tert</u> -butyl;  |
| for (2-8C)alkenyl:                             | vinyl, isopropenyl, allyl and but-2-enyl;   |
| for (2-8C)alkynyl:                             | ethynyl, 2-propynyl and but-2-ynyl;   |
| 10 for (1-6C)alkoxy:                           | methoxy, ethoxy, propoxy, isopropoxy and butoxy;  |
| for (2-6C)alkenyloxy:                          | vinylloxy and allyloxy;   |
| for (2-6C)alkynyloxy:                          | ethynyloxy and 2-propynyloxy;   |
| for (1-6C)alkylthio:                           | methylthio, ethylthio and propylthio;   |
| for (1-6C)alkylsulphinyl:                      | methylsulphinyl and ethylsulphinyl;   |
| 15 for (1-6C)alkylsulphonyl:                   | methylsulphonyl and ethylsulphonyl;   |
| for (1-6C)alkylamino:                          | methylamino, ethylamino, propylamino, isopropylamino and butylamino;  |
| for di-[(1-6C)alkyl]amino:                     | dimethylamino, diethylamino, <u>N</u> -ethyl- <u>N</u> -methylamino and diisopropylamino;                   |
| 20 for (1-6C)alkoxycarbonyl:                   | methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl and <u>tert</u> -butoxycarbonyl;                           |
| for <u>N</u> -(1-6C)alkylcarbamoyl:            | <u>N</u> -methylcarbamoyl, <u>N</u> -ethylcarbamoyl and <u>N</u> -propylcarbamoyl;                          |
| for <u>N,N</u> -di-[(1-6C)alkyl]carbamoyl:     | <u>N,N</u> -dimethylcarbamoyl, <u>N</u> -ethyl- <u>N</u> -methylcarbamoyl and <u>N,N</u> -diethylcarbamoyl; |
| 25 for (2-6C)alkanoyl:                         | acetyl and propionyl;   |
| for (2-6C)alkanoyloxy:                         | acetoxyl and propionyloxy;  |
| for (2-6C)alkanoylamino:                       | acetamido and propionamido;   |
| for <u>N</u> -(1-6C)alkyl-(2-6C)alkanoylamino: | <u>N</u> -methylacetamido and <u>N</u> -methylpropionamido;   |
| 30 for <u>N</u> -(1-6C)alkylsulphamoyl:        | <u>N</u> -methylsulphamoyl and <u>N</u> -ethylsulphamoyl;   |
| for <u>N,N</u> -di-[(1-6C)alkyl]sulphamoyl:    | <u>N,N</u> -dimethylsulphamoyl;   |
| for (1-6C)alkanesulphonylamino:                | methanesulphonylamino and ethanesulphonylamino;   |



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- for N-(1-6C)alkyl-(1-6C)alkanesulphonylamino: N-methylmethanesulphonylamino and  
N-methylethanesulphonylamino;
- for amino-(1-6C)alkyl: aminomethyl, 2-aminoethyl, 1-aminoethyl and  
3-aminopropyl;
- 5 for (1-6C)alkylamino-(1-6C)alkyl: methylaminomethyl, ethylaminomethyl,  
1-methylaminoethyl, 2-methylaminoethyl,  
2-ethylaminoethyl and 3-methylaminopropyl;
- for di-[(1-6C)alkyl]amino-(1-6C)alkyl: dimethylaminomethyl, diethylaminomethyl,  
1-dimethylaminoethyl, 2-dimethylaminoethyl and  
10 3-dimethylaminopropyl;
- for halogeno-(1-6C)alkyl: chloromethyl, 2-chloroethyl, 1-chloroethyl and  
3-chloropropyl;
- for hydroxy-(1-6C)alkyl: hydroxymethyl, 2-hydroxyethyl, 1-hydroxyethyl and  
3-hydroxypropyl;
- 15 for (1-6C)alkoxy-(1-6C)alkyl: methoxymethyl, ethoxymethyl, 1-methoxyethyl,  
2-methoxyethyl, 2-ethoxyethyl and  
3-methoxypropyl;
- for cyano-(1-6C)alkyl: cyanomethyl, 2-cyanoethyl, 1-cyanoethyl and  
3-cyanopropyl;
- 20 for (2-6C)alkanoylamino-(1-6C)alkyl: acetamidomethyl, propionamidomethyl and  
2-acetamidoethyl;
- for (1-6C)alkoxycarbonylamino-(1-6C)alkyl: methoxycarbonylaminomethyl,  
ethoxycarbonylaminomethyl,  
tert-butoxycarbonylaminomethyl and  
25 2-methoxycarbonylaminoethyl;
- for amino-(3-6C)alkyl: 3-aminopropyl;
- for (1-6C)alkylamino-(3-6C)alkyl: 3-methylaminopropyl;
- for di-[(1-6C)alkyl]amino-(3-6C)alkyl: 3-dimethylaminopropyl, 3-(N-ethyl-  
N-methylamino)propyl and 3-(N-isopropyl-  
30 N-methylamino)propyl;
- for hydroxy-(3-6C)alkyl: 3-hydroxypropyl;
- for (1-6C)alkoxy-(3-6C)alkyl: 3-methoxypropyl;
- for (2-6C)alkanoylamino-(3-6C)alkyl: 3-acetamidopropyl; and

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for (1-6C)alkoxycarbonylamino-(3-6C)alkyl: 3-methoxycarbonylaminoethyl.

When, as defined hereinbefore, an  $R^1$  or  $R^2$  group forms a group of the formula  $Q^1-X^1$  and, for example,  $X^1$  is a  $OC(R^4)_2$  linking group, it is the carbon atom, not the oxygen atom, of the  $OC(R^4)_2$  linking group which is attached to the quinazoline ring and the oxygen atom is  
 5 attached to the  $Q^1$  group. Similarly, when, for example a  $CH_3$  group within a  $R^1$  or  $R^2$  substituent bears a group of the formula  $-X^4-Q^2$  and, for example,  $X^4$  is a  $C(R^8)_2O$  linking group, it is the carbon atom, not the oxygen atom, of the  $C(R^8)_2O$  linking group which is attached to the  $CH_3$  group and the oxygen atom is linked to the  $Q^2$  group.

As defined hereinbefore, adjacent carbon atoms in any (2-6C)alkylene chain within a  
 10  $R^1$  or  $R^2$  substituent may be optionally separated by the insertion into the chain of a group such as O,  $CON(R^7)$  or  $C\equiv C$ . For example, insertion of a  $C\equiv C$  group into the ethylene chain within a 2-morpholinoethoxy group gives rise to a 4-morpholinobut-2-ynyloxy group and, for example, insertion of a CONH group into the ethylene chain within a 3-methoxypropoxy group gives rise to, for example, a 2-(2-methoxyacetamido)ethoxy group.

15 When, as defined hereinbefore, any  $CH_2$  or  $CH_3$  group within a  $R^1$  or  $R^2$  substituent optionally bears on each said  $CH_2$  or  $CH_3$  group one or more halogeno or (1-6C)alkyl substituents, there are suitably 1 or 2 halogeno or (1-6C)alkyl substituents present on each said  $CH_2$  group and there are suitably 1, 2 or 3 such substituents present on each said  $CH_3$  group.

When, as defined hereinbefore, any  $CH_2$  or  $CH_3$  group within a  $R^1$  or  $R^2$  substituent  
 20 optionally bears on each said  $CH_2$  or  $CH_3$  group a substituent as defined hereinbefore, suitable  $R^1$  or  $R^2$  substituents so formed include, for example, hydroxy-substituted heterocycl-  
 (1-6C)alkoxy groups such as 2-hydroxy-3-piperidinopropoxy and 2-hydroxy-  
 3-morpholinopropoxy, hydroxy-substituted amino-(2-6C)alkoxy groups such as 3-amino-  
 2-hydroxypropoxy, hydroxy-substituted (1-6C)alkylamino-(2-6C)alkoxy groups such as  
 25 2-hydroxy-3-methylaminopropoxy, hydroxy-substituted di-[(1-6C)alkyl]amino-(2-6C)alkoxy  
 groups such as 3-dimethylamino-2-hydroxypropoxy, hydroxy-substituted heterocycl-  
 (1-6C)alkylamino groups such as 2-hydroxy-3-piperidinopropylamino and 2-hydroxy-  
 3-morpholinopropylamino, hydroxy-substituted amino-(2-6C)alkylamino groups such as  
 3-amino-2-hydroxypropylamino, hydroxy-substituted (1-6C)alkylamino-(2-6C)alkylamino  
 30 groups such as 2-hydroxy-3-methylaminopropylamino, hydroxy-substituted  
 di-[(1-6C)alkyl]amino-(2-6C)alkylamino groups such as 3-dimethylamino-  
 2-hydroxypropylamino, hydroxy-substituted (1-6C)alkoxy groups such as 2-hydroxyethoxy,

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(1-6C)alkoxy-substituted (1-6C)alkoxy groups such as 2-methoxyethoxy and 3-ethoxypropoxy, (1-6C)alkylsulphonyl-substituted (1-6C)alkoxy groups such as 2-methylsulphonylethoxy and heterocyclyl-substituted (1-6C)alkylamino-(1-6C)alkyl groups such as 2-morpholinoethylaminomethyl, 2-piperazin-1-ylethylaminomethyl and

5 3-morpholinopropylaminomethyl.

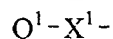
A suitable pharmaceutically-acceptable salt of a compound of the Formula I is, for example, an acid-addition salt of a compound of the Formula I, for example an acid-addition salt with an inorganic or organic acid such as hydrochloric, hydrobromic, sulphuric, trifluoroacetic, citric or maleic acid; or, for example, a salt of a compound of the Formula I

10 which is sufficiently acidic, for example an alkali or alkaline earth metal salt such as a calcium or magnesium salt, or an ammonium salt, or a salt with an organic base such as methylamine, dimethylamine, trimethylamine, piperidine, morpholine or tris-(2-hydroxyethyl)amine.

Particular novel compounds of the invention include, for example, quinazoline

15 derivatives of the Formula I, or pharmaceutically-acceptable salts thereof, wherein, unless otherwise stated, each of  $R^1$ ,  $R^2$  and  $R^3$  has any of the meanings defined hereinbefore or in paragraphs (a) to (j) hereinafter :-

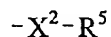
(a)  $R^1$  is hydrogen or methoxy and  $R^2$  is a group of the formula :



20 wherein  $X^1$  is selected from O,  $N(R^4)$ ,  $CON(R^4)$ ,  $N(R^4)CO$  and  $OC(R^4)_2$  wherein  $R^4$  is hydrogen or (1-6C)alkyl, and  $Q^1$  is heteroaryl, heteroaryl-(1-6C)alkyl, heteroaryloxy-(1-6C)alkyl, heterocyclyl, heterocyclyl-(1-6C)alkyl or heterocyclyloxy-(1-6C)alkyl, provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a piperidino-(2-4C)alkyl, morpholino-(2-4C)alkyl or piperazin-1-yl-(2-4C)alkyl group that is unsubstituted on the piperidino,

25 morpholino or piperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is excluded, and provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a 4-(1-4C)alkylpiperazin-1-yl-(2-4C)alkyl group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is also excluded,

30 or  $R^2$  is a group of the formula :



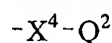
wherein  $X^2$  is selected from O and  $N(R^6)$ , wherein  $R^6$  is hydrogen or (1-6C)alkyl, and  $R^5$  is hydroxy-(3-6C)alkyl, (1-6C)alkoxy-(3-6C)alkyl, amino-(3-6C)alkyl, (1-6C)alkylamino-

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(3-6C)alkyl or di-[(1-6C)alkyl]amino-(3-6C)alkyl, provided that the central CH<sub>2</sub> group within any (CH<sub>2</sub>)<sub>3</sub> group within the (3-6C)alkylene group in a R<sup>5</sup> group bears a hydroxy or (2-6C)alkanoyloxy substituent,

and wherein adjacent carbon atoms in any (2-6C)alkylene chain within a R<sup>2</sup> substituent  
 5 are optionally separated by the insertion into the chain of a group selected from O, N(R<sup>5</sup>), CON(R<sup>5</sup>), N(R<sup>5</sup>)CO, CH=CH and C≡C wherein R<sup>5</sup> is hydrogen or (1-6C)alkyl,

and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a R<sup>2</sup> substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, cyano, amino, (1-6C)alkyl, (2-8C)alkenyl, (2-8C)alkynyl, (1-6C)alkoxy, (1-6C)alkylsulphonyl, (1-6C)alkylamino,  
 10 di-[(1-6C)alkyl]amino and (2-6C)alkanoyloxy, or from a group of the formula :

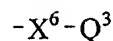


wherein X<sup>4</sup> is a direct bond or is selected from O, N(R<sup>8</sup>), CON(R<sup>8</sup>), N(R<sup>8</sup>)CO and C(R<sup>8</sup>)<sub>2</sub>O, wherein R<sup>8</sup> is hydrogen or (1-6C)alkyl, and Q<sup>2</sup> is heteroaryl, heteroaryl-(1-6C)alkyl, heterocyclyl or heterocyclyl-(1-6C)alkyl,

15 and wherein any aryl, heteroaryl or heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1, 2 or 3 substituents, which may be the same or different, selected from halogeno, trifluoromethyl, cyano, hydroxy, amino, carbamoyl, (1-6C)alkyl, (1-6C)alkoxy, (1-6C)alkoxycarbonyl, N-(1-6C)alkylcarbamoyl, N,N-di-[(1-6C)alkyl]carbamoyl and (2-6C)alkanoyl, or optionally bears 1 substituent selected from a group of the formula :



wherein X<sup>5</sup> is a direct bond or is selected from O and N(R<sup>10</sup>), wherein R<sup>10</sup> is hydrogen or (1-6C)alkyl, and R<sup>9</sup> is hydroxy-(1-6C)alkyl, (1-6C)alkoxy-(1-6C)alkyl, cyano-(1-6C)alkyl, amino-(1-6C)alkyl, (1-6C)alkylamino-(1-6C)alkyl, di-[(1-6C)alkyl]amino-(1-6C)alkyl, (2-6C)alkanoylamino-(1-6C)alkyl or (1-6C)alkoxycarbonylamino-(1-6C)alkyl, and from a  
 25 group of the formula :

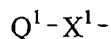


wherein X<sup>6</sup> is a direct bond or is selected from O, CO and N(R<sup>11</sup>), wherein R<sup>11</sup> is hydrogen or (1-6C)alkyl, and Q<sup>3</sup> is heterocyclyl or heterocyclyl-(1-6C)alkyl which optionally bears 1 or 2 substituents, which may be the same or different, selected from halogeno, (1-6C)alkyl and  
 30 (1-6C)alkoxy,

and wherein any heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 oxo substituents;

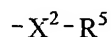
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(b)  $R^2$  is hydrogen or methoxy and  $R^1$  is a group of the formula :



wherein  $X^1$  is selected from O,  $N(R^4)$ ,  $CON(R^4)$ ,  $N(R^4)CO$  and  $OC(R^4)_2$  wherein  $R^4$  is hydrogen or (1-6C)alkyl, and  $Q^1$  is heteroaryl, heteroaryl-(1-6C)alkyl, heteroaryloxy-  
 5 (1-6C)alkyl, heterocyclyl, heterocyclyl-(1-6C)alkyl or heterocyclyloxy-(1-6C)alkyl, provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a piperidino-(2-4C)alkyl, morpholino-(2-4C)alkyl or piperazin-1-yl-(2-4C)alkyl group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is excluded, and provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a  
 10 4-(1-4C)alkylpiperazin-1-yl-(2-4C)alkyl group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is also excluded,

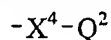
or  $R^1$  is a group of the formula :



15 wherein  $X^2$  is selected from O and  $N(R^6)$ , wherein  $R^6$  is hydrogen or (1-6C)alkyl, and  $R^5$  is hydroxy-(3-6C)alkyl, (1-6C)alkoxy-(3-6C)alkyl, amino-(3-6C)alkyl, (1-6C)alkylamino-(3-6C)alkyl or di-[(1-6C)alkyl]amino-(3-6C)alkyl, provided that the central  $CH_2$  group within any  $(CH_2)_3$  group within the (3-6C)alkylene group in a  $R^5$  group bears a hydroxy or (2-6C)alkanoyloxy substituent,

20 and wherein adjacent carbon atoms in any (2-6C)alkylene chain within a  $R^1$  substituent are optionally separated by the insertion into the chain of a group selected from O,  $N(R^5)$ ,  $CON(R^5)$ ,  $N(R^5)CO$ ,  $CH=CH$  and  $C\equiv C$  wherein  $R^5$  is hydrogen or (1-6C)alkyl,

and wherein any  $CH_2$  or  $CH_3$  group within a  $R^1$  substituent optionally bears on each said  $CH_2$  or  $CH_3$  group a substituent selected from hydroxy, cyano, amino, (1-6C)alkyl,  
 25 (2-8C)alkenyl, (2-8C)alkynyl, (1-6C)alkoxy, (1-6C)alkylsulphonyl, (1-6C)alkylamino, di-[(1-6C)alkyl]amino and (2-6C)alkanoyloxy, or from a group of the formula :

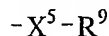


wherein  $X^4$  is a direct bond or is selected from O,  $N(R^8)$ ,  $CON(R^8)$ ,  $N(R^8)CO$  and  $C(R^8)_2O$ , wherein  $R^8$  is hydrogen or (1-6C)alkyl, and  $Q^2$  is heteroaryl, heteroaryl-(1-6C)alkyl,  
 30 heterocyclyl or heterocyclyl-(1-6C)alkyl,

and wherein any aryl, heteroaryl or heterocyclyl group within a substituent on  $R^1$  optionally bears 1, 2 or 3 substituents, which may be the same or different, selected from

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halogeno, trifluoromethyl, cyano, hydroxy, amino, carbamoyl, (1-6C)alkyl, (1-6C)alkoxy, (1-6C)alkoxycarbonyl, N-(1-6C)alkylcarbamoyl, N,N-di-[(1-6C)alkyl]carbamoyl and (2-6C)alkanoyl, or optionally bears 1 substituent selected from a group of the formula :



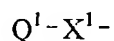
- 5 wherein  $X^5$  is a direct bond or is selected from O and  $N(R^{10})$ , wherein  $R^{10}$  is hydrogen or (1-6C)alkyl, and  $R^9$  is hydroxy-(1-6C)alkyl, (1-6C)alkoxy-(1-6C)alkyl, cyano-(1-6C)alkyl, amino-(1-6C)alkyl, (1-6C)alkylamino-(1-6C)alkyl, di-[(1-6C)alkyl]amino-(1-6C)alkyl, (2-6C)alkanoylamino-(1-6C)alkyl or (1-6C)alkoxycarbonylamino-(1-6C)alkyl, and from a group of the formula :



wherein  $X^6$  is a direct bond or is selected from O, CO and  $N(R^{11})$ , wherein  $R^{11}$  is hydrogen or (1-6C)alkyl, and  $Q^3$  is heterocyclyl or heterocyclyl-(1-6C)alkyl which optionally bears 1 or 2 substituents, which may be the same or different, selected from halogeno, (1-6C)alkyl and (1-6C)alkoxy,

- 15 and wherein any heterocyclyl group within a substituent on  $R^1$  optionally bears 1 or 2 oxo substituents;

(c)  $R^1$  is hydrogen or methoxy and  $R^2$  is a group of the formula :



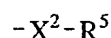
wherein  $X^1$  is selected from O, NH, CONH, NHCO and  $OCH_2$  and  $Q^1$  is 2-thienyl,

- 20 1-imidazolyl, 1,2,3-triazol-1-yl, 1,2,4-triazol-1-yl, 2-, 3- or 4-pyridyl, 2-imidazol-1-ylethyl, 3-imidazol-1-ylpropyl, 2-(1,2,3-triazolyl)ethyl, 3-(1,2,3-triazolyl)propyl, 2-(1,2,4-triazolyl)ethyl, 3-(1,2,4-triazolyl)propyl, 2-, 3- or 4-pyridylmethyl, 2-(2-, 3- or 4-pyridyl)ethyl, 3-(2-, 3- or 4-pyridyl)propyl, 2-(2-, 3- or 4-pyridyloxy)ethyl, 3-(2-, 3- or 4-pyridyloxy)propyl, 1-, 2- or 3-pyrrolidinyl, morpholino, 1,1-dioxotetrahydro-  
 25 4H-1,4-thiazin-4-yl, piperidino, piperidin-3-yl, piperidin-4-yl, 1-, 3- or 4-homopiperidinyl, piperazin-1-yl, homopiperazin-1-yl, 1-, 2- or 3-pyrrolidinylmethyl, morpholinomethyl, piperidinomethyl, 3- or 4-piperidinylmethyl, 1-, 3- or 4-homopiperidinylmethyl, 2-azetidin-1-ylethyl, 3-azetidin-1-ylpropyl, 2-pyrrolidin-1-ylethyl, 3-pyrrolidin-2-ylpropyl, pyrrolidin-2-ylmethyl, 2-pyrrolidin-2-ylethyl, 3-pyrrolidin-1-ylpropyl, 2-morpholinoethyl,  
 30 3-morpholinopropyl, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)ethyl, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propyl, 2-piperidinoethyl, 3-piperidinopropyl, 2-piperidin-3-ylethyl, 2-piperidin-4-ylethyl, 2-homopiperidin-1-ylethyl, 3-homopiperidin-1-ylpropyl, 2-piperazin-1-ylethyl, 3-piperazin-1-ylpropyl, 2-homopiperazin-1-ylethyl or

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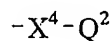
3-homopiperazin-1-ylpropyl, provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a piperidino-(2-4C)alkyl, morpholino-(2-4C)alkyl or piperazin-1-yl-(2-4C)alkyl group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is excluded, and provided that, when  $X^1$  is O, any compound wherein  
 5  $Q^1$  is a 4-(1-4C)alkylpiperazin-1-yl-(2-4C)alkyl group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is also excluded,

or  $R^2$  is a group of the formula :



10 wherein  $X^2$  is selected from O and NH and  $R^5$  is 3-hydroxypropyl, 3-methoxypropyl, 3-ethoxypropyl, 3-aminopropyl, 3-methylaminopropyl, 3-ethylaminopropyl, 3-isopropylaminopropyl, 3-dimethylaminopropyl, 3-diethylaminopropyl, 3-(N-cyclobutyl-N-methylamino)propyl, 3-(N-ethyl-N-methylamino)propyl, 3-(N-ethyl-N-isopropylamino)propyl or 3-(N-isopropyl-N-methylamino)propyl, provided that  
 15 the central  $CH_2$  group within the propyl group bears a hydroxy or acetoxy substituent, and wherein adjacent carbon atoms in any (2-6C)alkylene chain within a  $R^2$  substituent are optionally separated by the insertion into the chain of a group selected from O, NH, CONH, NHCO,  $CH=CH$  and  $C\equiv C$ ,

and wherein any  $CH_2$  or  $CH_3$  group within a  $R^2$  substituent optionally bears on each  
 20 said  $CH_2$  or  $CH_3$  group a substituent selected from hydroxy, cyano, amino, methyl, ethyl, vinyl, ethynyl, methoxy, methylsulphonyl, methylamino, dimethylamino, acetoxy, propionyloxy, butyryloxy, isobutyryloxy, isopentanoyloxy, cyclobutylcarbonyl, cyclopentylcarbonyl and cyclohexylcarbonyl, or from a group of the formula :

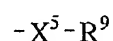


25 wherein  $X^4$  is a direct bond or is selected from O, NH, CONH, NHCO and  $CH_2O$  and  $Q^2$  is pyridyl, pyridylmethyl, pyrrolidin-1-yl, pyrrolidin-2-yl, morpholino, piperidino, piperidin-3-yl, piperidin-4-yl, piperazin-1-yl, 2-pyrrolidin-1-ylethyl, 3-pyrrolidin-1-ylpropyl, pyrrolidin-2-ylmethyl, 2-pyrrolidin-2-ylethyl, 3-pyrrolidin-2-ylpropyl, 2-morpholinoethyl, 3-morpholinopropyl, 2-piperidinoethyl, 3-piperidinopropyl, piperidin-3-ylmethyl, 2-piperidin-3-ylethyl, piperidin-4-ylmethyl, 2-piperidin-4-ylethyl, 2-piperazin-1-ylethyl or 3-piperazin-1-ylpropyl,  
 30

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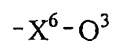
and wherein any aryl, heteroaryl or heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1, 2 or 3 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, cyano, hydroxy, amino, carbamoyl, methyl, ethyl, cyclopropyl, allyl, methoxy, acetyl and tert-butoxycarbonyl, or optionally bears 1 substituent selected from

5 a group of the formula :



wherein X<sup>5</sup> is a direct bond or is selected from O and NH and R<sup>9</sup> is 2-hydroxyethyl, 3-hydroxypropyl, 2-methoxyethyl, 3-methoxypropyl, cyanomethyl, aminomethyl, 2-aminoethyl, 3-aminopropyl, methylaminomethyl, 2-methylaminoethyl,

10 3-methylaminopropyl, 2-ethylaminoethyl, 3-ethylaminopropyl, dimethylaminomethyl, 2-dimethylaminoethyl, 3-dimethylaminopropyl, acetamidomethyl, methoxycarbonylaminomethyl, ethoxycarbonylaminomethyl or tert-butoxycarbonylaminomethyl, and from a group of the formula :

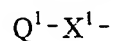


15 wherein X<sup>6</sup> is a direct bond or is selected from O and NH and Q<sup>3</sup> is pyrrolidin-1-ylmethyl, 2-pyrrolidin-1-ylethyl, 3-pyrrolidin-1-ylpropyl, morpholinomethyl, 2-morpholinoethyl, 3-morpholinopropyl, piperidinomethyl, 2-piperidinoethyl, 3-piperidinopropyl, piperazin-1-ylmethyl, 2-piperazin-1-ylethyl or 3-piperazin-1-ylpropyl, each of which optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro,

20 chloro, methyl and methoxy,

and wherein any heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 oxo substituents;

(d) R<sup>2</sup> is hydrogen or methoxy and R<sup>1</sup> is a group of the formula :



25 wherein X<sup>1</sup> is selected from O, NH, CONH, NHCO and OCH<sub>2</sub> and Q<sup>1</sup> is 2-thienyl, 1-imidazolyl, 1,2,3-triazol-1-yl, 1,2,4-triazol-1-yl, 2-, 3- or 4-pyridyl, 2-imidazol-1-ylethyl, 3-imidazol-1-ylpropyl, 2-(1,2,3-triazolyl)ethyl, 3-(1,2,3-triazolyl)propyl, 2-(1,2,4-triazolyl)ethyl, 3-(1,2,4-triazolyl)propyl, 2-, 3- or 4-pyridylmethyl, 2-(2-, 3- or 4-pyridyl)ethyl, 3-(2-, 3- or 4-pyridyl)propyl, 2-(2-, 3- or 4-pyridyloxy)ethyl,

30 3-(2-, 3- or 4-pyridyloxy)propyl, 1-, 2- or 3-pyrrolidinyl, morpholino, 1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl, piperidino, piperidin-3-yl, piperidin-4-yl, 1-, 3- or 4-homopiperidinyl, piperazin-1-yl, homopiperazin-1-yl, 1-, 2- or 3-pyrrolidinylmethyl, morpholinomethyl, piperidinomethyl, 3- or 4-piperidinylmethyl, 1-, 3- or 4-homopiperidinylmethyl,



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2-azetidin-1-ylethyl, 3-azetidin-1-ylpropyl, 2-pyrrolidin-1-ylethyl, 3-pyrrolidin-2-ylpropyl, pyrrolidin-2-ylmethyl, 2-pyrrolidin-2-ylethyl, 3-pyrrolidin-1-ylpropyl, 2-morpholinoethyl, 3-morpholinopropyl, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)ethyl, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propyl, 2-piperidinoethyl, 3-piperidinopropyl, 2-piperidin-3-ylethyl, 2-piperidin-4-ylethyl, 2-homopiperidin-1-ylethyl, 3-homopiperidin-1-ylpropyl, 2-piperazin-1-ylethyl, 3-piperazin-1-ylpropyl, 2-homopiperazin-1-ylethyl or 3-homopiperazin-1-ylpropyl, provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a piperidino-(2-4C)alkyl, morpholino-(2-4C)alkyl or piperazin-1-yl-(2-4C)alkyl group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is excluded, and provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a 4-(1-4C)alkylpiperazin-1-yl-(2-4C)alkyl group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is also excluded,

or  $R^1$  is a group of the formula :

15  $-X^2-R^5$

wherein  $X^2$  is selected from O and NH and  $R^5$  is 3-hydroxypropyl, 3-methoxypropyl, 3-ethoxypropyl, 3-aminopropyl, 3-methylaminopropyl, 3-ethylaminopropyl, 3-isopropylaminopropyl, 3-dimethylaminopropyl, 3-diethylaminopropyl, 3-(N-cyclobutyl-N-methylamino)propyl, 3-(N-ethyl-N-methylamino)propyl, 3-(N-ethyl-N-isopropylamino)propyl or 3-(N-isopropyl-N-methylamino)propyl, provided that the central  $CH_2$  group within the propyl group bears a hydroxy or acetoxy substituent,

and wherein adjacent carbon atoms in any (2-6C)alkylene chain within a  $R^1$  substituent are optionally separated by the insertion into the chain of a group selected from O, NH, CONH, NHCO,  $CH=CH$  and  $C\equiv C$ ,

25 and wherein any  $CH_2$  or  $CH_3$  group within a  $R^1$  substituent optionally bears on each said  $CH_2$  or  $CH_3$  group a substituent selected from hydroxy, cyano, amino, methyl, ethyl, vinyl, ethynyl, methoxy, methylsulphonyl, methylamino, dimethylamino, acetoxy, propionyloxy, butyryloxy, isobutyryloxy, isopentanoyloxy, cyclobutylcarbonyl, cyclopentylcarbonyl and cyclohexylcarbonyl, or from a group of the formula :

30  $-X^4-Q^2$

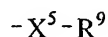
wherein  $X^4$  is a direct bond or is selected from O, NH, CONH, NHCO and  $CH_2O$  and  $Q^2$  is pyridyl, pyridylmethyl, pyrrolidin-1-yl, pyrrolidin-2-yl, morpholino, piperidino, piperidin-3-yl,

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piperidin-4-yl, piperazin-1-yl, 2-pyrrolidin-1-ylethyl, 3-pyrrolidin-1-ylpropyl, pyrrolidin-2-ylmethyl, 2-pyrrolidin-2-ylethyl, 3-pyrrolidin-2-ylpropyl, 2-morpholinoethyl, 3-morpholinopropyl, 2-piperidinoethyl, 3-piperidinopropyl, piperidin-3-ylmethyl, 2-piperidin-3-ylethyl, piperidin-4-ylmethyl, 2-piperidin-4-ylethyl, 2-piperazin-1-ylethyl or 3-piperazin-1-ylpropyl,

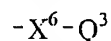
and wherein any aryl, heteroaryl or heterocyclyl group within a substituent on R<sup>1</sup> optionally bears 1, 2 or 3 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, cyano, hydroxy, amino, carbamoyl, methyl, ethyl, cyclopropyl, allyl, methoxy, acetyl and tert-butoxycarbonyl, or optionally bears 1 substituent selected from

10 a group of the formula :



wherein X<sup>5</sup> is a direct bond or is selected from O and NH and R<sup>9</sup> is 2-hydroxyethyl, 3-hydroxypropyl, 2-methoxyethyl, 3-methoxypropyl, cyanomethyl, aminomethyl, 2-aminoethyl, 3-aminopropyl, methylaminomethyl, 2-methylaminoethyl,

15 3-methylaminopropyl, 2-ethylaminoethyl, 3-ethylaminopropyl, dimethylaminomethyl, 2-dimethylaminoethyl, 3-dimethylaminopropyl, acetamidomethyl, methoxycarbonylaminomethyl, ethoxycarbonylaminomethyl or tert-butoxycarbonylaminomethyl, and from a group of the formula :



20 wherein X<sup>6</sup> is a direct bond or is selected from O and NH and Q<sup>3</sup> is pyrrolidin-1-ylmethyl, 2-pyrrolidin-1-ylethyl, 3-pyrrolidin-1-ylpropyl, morpholinomethyl, 2-morpholinoethyl, 3-morpholinopropyl, piperidinomethyl, 2-piperidinoethyl, 3-piperidinopropyl, piperazin-1-ylmethyl, 2-piperazin-1-ylethyl or 3-piperazin-1-ylpropyl, each of which optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro,

25 chloro, methyl and methoxy,

and wherein any heterocyclyl group within a substituent on R<sup>1</sup> optionally bears 1 or 2 oxo substituents;

(e) R<sup>1</sup> is hydrogen or methoxy and R<sup>2</sup> is 2-imidazol-1-ylethoxy, 3-imidazol-1-ylpropoxy, 2-(1,2,3-triazol-1-yl)ethoxy, 3-(1,2,3-triazol-1-yl)propoxy,

30 pyrid-2-ylmethoxy, pyrid-3-ylmethoxy, pyrid-4-ylmethoxy, 2-pyrid-2-ylethoxy, 2-pyrid-3-ylethoxy, 2-pyrid-4-ylethoxy, 3-pyrid-2-ylpropoxy, 3-pyrid-3-ylpropoxy, 3-pyrid-4-ylpropoxy, 2-pyrid-2-yloxyethoxy, 2-pyrid-3-yloxyethoxy, 2-pyrid-4-yloxyethoxy, 3-pyrid-2-yloxypropoxy, 3-pyrid-3-yloxypropoxy, 3-pyrid-4-yloxypropoxy, pyrrolidin-1-yl,

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- morpholino, piperidino, piperazin-1-yl, 2-pyrrolidin-1-ylethoxy, 3-pyrrolidin-1-ylpropoxy, pyrrolidin-3-yloxy, pyrrolidin-2-ylmethoxy, 2-pyrrolidin-2-ylethoxy, 3-pyrrolidin-2-ylpropoxy, 2-morpholinoethoxy, 3-morpholinopropoxy, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)ethoxy, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy,
- 5 2-azetidin-1-ylethoxy, 3-azetidin-1-ylpropoxy, 2-piperidinoethoxy, 3-piperidinopropoxy, piperidin-3-yloxy, piperidin-4-yloxy, piperidin-3-ylmethoxy, 2-piperidin-3-ylethoxy, piperidin-4-ylmethoxy, 2-piperidin-4-ylethoxy, 2-homopiperidin-1-ylethoxy, 3-homopiperidin-1-ylpropoxy, 2-piperazin-1-ylethoxy, 3-piperazin-1-ylpropoxy, 2-homopiperazin-1-ylethoxy, 3-homopiperazin-1-ylpropoxy, 2-pyrrolidin-1-ylethylamino,
- 10 3-pyrrolidin-1-ylpropylamino, pyrrolidin-3-ylamino, pyrrolidin-2-ylmethylamino, 2-pyrrolidin-2-ylethylamino, 3-pyrrolidin-2-ylpropylamino, 2-morpholinoethylamino, 3-morpholinopropylamino, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)ethylamino, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propylamino, 2-piperidinoethylamino, 3-piperidinopropylamino, piperidin-3-ylamino, piperidin-4-ylamino,
- 15 piperidin-3-ylmethylamino, 2-piperidin-3-ylethylamino, piperidin-4-ylmethylamino, 2-piperidin-4-ylethylamino, 2-homopiperidin-1-ylethylamino, 3-homopiperidin-1-ylpropylamino, 2-piperazin-1-ylethylamino, 3-piperazin-1-ylpropylamino, 2-homopiperazin-1-ylethylamino or 3-homopiperazin-1-ylpropylamino, provided that, any compound wherein  $R^2$  is a piperidino-(2-4C)alkoxy, morpholino-(2-4C)alkoxy or piperazin-1-yl-(2-4C)alkoxy
- 20 group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the (2-4C)alkyleneoxy group is excluded, and provided that any compound wherein  $R^2$  is a 4-(1-4C)alkylpiperazin-1-yl-(2-4C)alkoxy group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the (2-4C)alkyleneoxy group is also excluded,
- 25 or  $R^2$  is a group selected from 3-hydroxypropoxy, 3-methoxypropoxy, 3-ethoxypropoxy, 3-aminopropoxy, 3-methylaminopropoxy, 3-ethylaminopropoxy, 3-isopropylaminopropoxy, 3-dimethylaminopropoxy, 3-diethylaminopropoxy, 3-(N-cyclobutyl-N-methylamino)propoxy, 3-(N-ethyl-N-methylamino)propoxy, 3-(N-ethyl-N-isopropylamino)propoxy or 3-(N-isopropyl-N-methylamino)propoxy, provided
- 30 that the central  $CH_2$  group within the propoxy group bears a hydroxy or acetoxy substituent,

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and wherein adjacent carbon atoms in any (2-6C)alkylene chain within a R<sup>2</sup> substituent are optionally separated by the insertion into the chain of a group selected from O, NH, CH=CH and C≡C,

- and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a R<sup>2</sup> substituent optionally bears on each
- 5 said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, cyano, amino, methyl, ethyl, vinyl, ethynyl, methoxy, methylsulphonyl, methylamino, dimethylamino and acetoxy,
- and wherein any pyridyl or heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, hydroxy, amino, carbamoyl, methyl, ethyl and methoxy, and a
- 10 piperidin-3-yl, piperidin-4-yl or piperazin-1-yl group within a R<sup>2</sup> substituent is optionally N-substituted with cyclopropyl, allyl, acetyl, 2-methoxyethyl, 3-methoxypropyl, cyanomethyl, 2-aminoethyl, 3-aminopropyl, 2-methylaminoethyl, 3-methylaminopropyl, 2-dimethylaminoethyl, 3-dimethylaminopropyl, 2-pyrrolidin-1-ylethyl, 3-pyrrolidin-1-ylpropyl, 2-morpholinoethyl, 3-morpholinopropyl, 2-piperidinoethyl,
- 15 3-piperidinopropyl, 2-piperazin-1-ylethyl or 3-piperazin-1-ylpropyl, the last 8 of which substituents each optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, methyl and methoxy,

and wherein any heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 oxo substituents;

- 20 (f) R<sup>2</sup> is hydrogen or methoxy and R<sup>1</sup> is 2-imidazol-1-ylethoxy, 3-imidazol-1-ylpropoxy, 2-(1,2,3-triazol-1-yl)ethoxy, 3-(1,2,3-triazol-1-yl)propoxy, pyrid-2-ylmethoxy, pyrid-3-ylmethoxy, pyrid-4-ylmethoxy, 2-pyrid-2-ylethoxy, 2-pyrid-3-ylethoxy, 2-pyrid-4-ylethoxy, 3-pyrid-2-ylpropoxy, 3-pyrid-3-ylpropoxy, 3-pyrid-4-ylpropoxy, 2-pyrid-2-yloxyethoxy, 2-pyrid-3-yloxyethoxy, 2-pyrid-4-yloxyethoxy,
- 25 3-pyrid-2-yloxypropoxy, 3-pyrid-3-yloxypropoxy, 3-pyrid-4-yloxypropoxy, pyrrolidin-1-yl, morpholino, piperidino, piperazin-1-yl, 2-azetidin-1-ylethoxy, 3-azetidin-1-ylpropoxy, 2-pyrrolidin-1-ylethoxy, 3-pyrrolidin-1-ylpropoxy, pyrrolidin-3-yloxy, pyrrolidin-2-ylmethoxy, 2-pyrrolidin-2-ylethoxy, 3-pyrrolidin-2-ylpropoxy, 2-morpholinoethoxy, 3-morpholinopropoxy, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-
- 30 4-yl)ethoxy, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-piperidinoethoxy, 3-piperidinopropoxy, piperidin-3-yloxy, piperidin-4-yloxy, piperidin-3-ylmethoxy, 2-piperidin-3-ylethoxy, piperidin-4-ylmethoxy, 2-piperidin-4-ylethoxy, 2-homopiperidin-

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- 1-ylethoxy, 3-homopiperidin-1-ylpropoxy, 2-piperazin-1-ylethoxy, 3-piperazin-1-ylpropoxy, 2-homopiperazin-1-ylethoxy, 3-homopiperazin-1-ylpropoxy, 2-pyrrolidin-1-ylethylamino, 3-pyrrolidin-1-ylpropylamino, pyrrolidin-3-ylamino, pyrrolidin-2-ylmethylamino, 2-pyrrolidin-2-ylethylamino, 3-pyrrolidin-2-ylpropylamino, 2-morpholinoethylamino, 3-morpholinopropylamino, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)ethylamino, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propylamino, 2-piperidinoethylamino, 3-piperidinopropylamino, piperidin-3-ylamino, piperidin-4-ylamino, piperidin-3-ylmethylamino, 2-piperidin-3-ylethylamino, piperidin-4-ylmethylamino, 2-piperidin-4-ylethylamino, 2-homopiperidin-1-ylethylamino, 3-homopiperidin-1-ylpropylamino, 2-piperazin-1-ylethylamino, 3-piperazin-1-ylpropylamino, 2-homopiperazin-1-ylethylamino or 3-homopiperazin-1-ylpropylamino, provided that, any compound wherein R<sup>1</sup> is a piperidino-(2-4C)alkoxy, morpholino-(2-4C)alkoxy or piperazin-1-yl-(2-4C)alkoxy group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the (2-4C)alkyleneoxy group is excluded, and provided that any compound wherein R<sup>1</sup> is a 4-(1-4C)alkylpiperazin-1-yl-(2-4C)alkoxy group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the (2-4C)alkyleneoxy group is also excluded,
- or R<sup>1</sup> is a group selected from 3-hydroxypropoxy, 3-methoxypropoxy, 3-ethoxypropoxy, 3-aminopropoxy, 3-methylaminopropoxy, 3-ethylaminopropoxy, 3-isopropylaminopropoxy, 3-dimethylaminopropoxy, 3-diethylaminopropoxy, 3-(N-cyclobutyl-N-methylamino)propoxy, 3-(N-ethyl-N-methylamino)propoxy, 3-(N-ethyl-N-isopropylamino)propoxy or 3-(N-isopropyl-N-methylamino)propoxy, provided that the central CH<sub>2</sub> group within the propoxy group bears a hydroxy or acetoxy substituent, and wherein adjacent carbon atoms in any (2-6C)alkylene chain within a R<sup>1</sup> substituent are optionally separated by the insertion into the chain of a group selected from O, NH, CH=CH and C≡C,
- and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a R<sup>1</sup> substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, cyano, amino, methyl, ethyl, vinyl, ethynyl, methoxy, methylsulphonyl, methylamino, dimethylamino and acetoxy, and wherein any pyridyl or heterocyclyl group within a substituent on R<sup>1</sup> optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, hydroxy, amino, carbamoyl, methyl, ethyl and methoxy, and a

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piperidin-3-yl, piperidin-4-yl or piperazin-1-yl group within a R<sup>1</sup> substituent is optionally N-substituted with cyclopropyl, allyl, acetyl, 2-methoxyethyl, 3-methoxypropyl, cyanomethyl, 2-aminoethyl, 3-aminopropyl, 2-methylaminoethyl, 3-methylaminopropyl, 2-dimethylaminoethyl, 3-dimethylaminopropyl, 2-pyrrolidin-1-ylethyl, 3-pyrrolidin-1-ylpropyl, 2-morpholinoethyl, 3-morpholinopropyl, 2-piperidinoethyl, 3-piperidinopropyl, 2-piperazin-1-ylethyl or 3-piperazin-1-ylpropyl, the last 8 of which substituents each optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, methyl and methoxy,

and wherein any heterocyclyl group within a substituent on R<sup>1</sup> optionally bears 1 or 2  
 10 oxo substituents;

- (g) R<sup>1</sup> is methoxy and R<sup>2</sup> is 2-imidazol-1-ylethoxy, 3-(1,2,3-triazol-1-yl)propoxy, 4-pyridylmethoxy, 2-pyrid-4-yloxyethoxy, 3-pyrid-2-yloxypropoxy, 3-pyrid-4-yloxypropoxy, 2-pyrrolidin-1-ylethoxy, 3-pyrrolidin-1-ylpropoxy, pyrrolidin-3-yloxy, pyrrolidin-2-ylmethoxy, 2-pyrrolidin-2-ylethoxy, 3-pyrrolidin-2-ylpropoxy, 2-morpholinoethoxy, 3-morpholinopropoxy, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)ethoxy, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-piperidinoethoxy, 3-piperidinopropoxy, piperidin-3-yloxy, piperidin-4-yloxy, piperidin-3-ylmethoxy, 2-piperidin-3-ylethoxy, piperidin-4-ylmethoxy, 2-piperidin-4-ylethoxy, 2-homopiperidin-1-ylethoxy, 3-homopiperidin-1-ylpropoxy, 2-piperazin-1-ylethoxy, 3-piperazin-1-ylpropoxy, 2-homopiperazin-1-ylethoxy or 3-homopiperazin-1-ylpropoxy, provided that, any compound wherein R<sup>2</sup> is a 2-piperidinoethoxy, 3-piperidinopropoxy, 2-morpholinoethoxy, 3-morpholinopropoxy, 2-piperazin-1-ylethoxy or 3-piperazin-1-ylpropoxy group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the ethoxy or propoxy group is excluded, and  
 25 provided that any compound wherein R<sup>2</sup> is a 2-[4-(1-4C)alkyl]piperazin-1-ylethoxy or 3-[4-(1-4C)alkyl]piperazin-1-ylpropoxy group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the ethoxy or propoxy group is also excluded,

or R<sup>2</sup> is a group selected from 3-aminopropoxy, 3-methylaminopropoxy, 3-ethylaminopropoxy, 3-isopropylaminopropoxy, 3-dimethylaminopropoxy, 3-diethylaminopropoxy or 3-(N-isopropyl-N-methylamino)propoxy, provided that the central CH<sub>2</sub> group within the propoxy group bears a hydroxy or acetoxy substituent,

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and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a R<sup>2</sup> substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, cyano, amino, methoxy, methylsulphonyl, methylamino and dimethylamino,

and wherein any pyridyl or heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, cyano, hydroxy, amino, methyl, ethyl and methoxy,

and wherein any heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 oxo substituents;

(h) R<sup>2</sup> is methoxy and R<sup>1</sup> is 2-imidazol-1-ylethoxy, 3-(1,2,3-triazol-1-yl)propoxy, 4-pyridylmethoxy, 2-pyrid-4-yloxyethoxy, 3-pyrid-2-yloxypropoxy, 3-pyrid-4-yloxypropoxy, 2-pyrrolidin-1-ylethoxy, 3-pyrrolidin-1-ylpropoxy, pyrrolidin-3-yloxy, pyrrolidin-2-ylmethoxy, 2-pyrrolidin-2-ylethoxy, 3-pyrrolidin-2-ylpropoxy, 2-morpholinoethoxy, 3-morpholinopropoxy, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)ethoxy, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-piperidinoethoxy, 3-piperidinopropoxy, piperidin-3-yloxy, piperidin-4-yloxy, piperidin-3-ylmethoxy, 2-piperidin-3-ylethoxy, piperidin-4-ylmethoxy, 2-piperidin-4-ylethoxy, 2-homopiperidin-1-ylethoxy, 3-homopiperidin-1-ylpropoxy, 2-piperazin-1-ylethoxy, 3-piperazin-1-ylpropoxy, 2-homopiperazin-1-ylethoxy or 3-homopiperazin-1-ylpropoxy, provided that, any compound wherein R<sup>1</sup> is a 2-piperidinoethoxy, 3-piperidinopropoxy, 2-morpholinoethoxy, 3-morpholinopropoxy, 2-piperazin-1-ylethoxy or 3-piperazin-1-ylpropoxy group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the ethoxy or propoxy group is excluded, and provided that any compound wherein R<sup>1</sup> is a 2-[4-(1-4C)alkyl]piperazin-1-ylethoxy or 3-[4-(1-4C)alkyl]piperazin-1-ylpropoxy group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the ethoxy or propoxy group is also excluded,

or R<sup>1</sup> is a group selected from 3-aminopropoxy, 3-methylaminopropoxy, 3-ethylaminopropoxy, 3-isopropylaminopropoxy, 3-dimethylaminopropoxy, 3-diethylaminopropoxy or 3-(N-isopropyl-N-methylamino)propoxy, provided that the central CH<sub>2</sub> group within the propoxy group bears a hydroxy or acetoxy substituent,

and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a R<sup>1</sup> substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, cyano, amino, methoxy, methylsulphonyl, methylamino and dimethylamino,

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and wherein any pyridyl or heterocyclyl group within a substituent on R<sup>1</sup> optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, cyano, hydroxy, amino, methyl, ethyl and methoxy,

and wherein any heterocyclyl group within a substituent on R<sup>1</sup> optionally bears 1 or 2  
5 oxo substituents;

- (i) R<sup>3</sup> is chloro or bromo; and
- (j) R<sup>3</sup> is chloro.

A particular compound of the invention is a quinazoline derivative of the Formula I wherein :

10 R<sup>1</sup> is methoxy and R<sup>2</sup> is 2-imidazol-1-ylethoxy, 3-(1,2,3-triazol-1-yl)propoxy, 4-pyridylmethoxy, 2-pyrid-4-yloxyethoxy, 3-pyrid-2-yloxypropoxy, 3-pyrid-4-yloxypropoxy, 2-pyrrolidin-1-ylethoxy, 3-pyrrolidin-1-ylpropoxy, pyrrolidin-3-yloxy, pyrrolidin-2-ylmethoxy, 2-pyrrolidin-2-ylethoxy, 3-pyrrolidin-2-ylpropoxy, 2-morpholinoethoxy, 3-morpholinopropoxy, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-  
15 4-yl)ethoxy, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-piperidinoethoxy, 3-piperidinopropoxy, piperidin-3-yloxy, piperidin-4-yloxy, piperidin-3-ylmethoxy, 2-piperidin-3-ylethoxy, piperidin-4-ylmethoxy, 2-piperidin-4-ylethoxy, 2-homopiperidin-1-ylethoxy, 3-homopiperidin-1-ylpropoxy, 2-piperazin-1-ylethoxy, 3-piperazin-1-ylpropoxy, 2-homopiperazin-1-ylethoxy or 3-homopiperazin-1-ylpropoxy,  
20 provided that, any compound wherein R<sup>2</sup> is a 2-piperidinoethoxy, 3-piperidinopropoxy, 2-morpholinoethoxy, 3-morpholinopropoxy, 2-piperazin-1-ylethoxy or 3-piperazin-1-ylpropoxy group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the ethoxy or propoxy group is excluded, and provided that any compound wherein R<sup>2</sup> is a 2-[4-(1-4C)alkyl]piperazin-1-ylethoxy or  
25 3-[4-(1-4C)alkyl]piperazin-1-ylpropoxy group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the ethoxy or propoxy group is also excluded,

or R<sup>2</sup> is a group selected from 3-aminopropoxy, 3-methylaminopropoxy, 3-ethylaminopropoxy, 3-isopropylaminopropoxy, 3-dimethylaminopropoxy,  
30 3-diethylaminopropoxy or 3-(N-isopropyl-N-methylamino)propoxy, provided that the central CH<sub>2</sub> group within the propoxy group bears a hydroxy or acetoxy substituent,



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and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a R<sup>2</sup> substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, cyano, amino, methoxy, methylsulphonyl, methylamino and dimethylamino,

and wherein any pyridyl or heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, cyano, hydroxy, amino, methyl, ethyl and methoxy,

and wherein any heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 oxo substituents; and

R<sup>3</sup> is chloro or bromo;

10 or a pharmaceutically-acceptable acid-addition salt thereof.

A further particular compound of the invention is a quinazoline derivative of the Formula I wherein :

R<sup>1</sup> is methoxy and R<sup>2</sup> is 2-imidazol-1-ylethoxy, 3-(1,2,3-triazol-1-yl)propoxy, 4-pyridylmethoxy, 2-pyrid-4-yloxyethoxy, 3-pyrid-2-yloxypropoxy, 3-pyrid-4-yloxypropoxy, 15 3-azetidin-1-yl-2-hydroxypropoxy, 2-pyrrolidin-1-ylethoxy, 3-pyrrolidin-1-ylpropoxy, 2-hydroxy-3-pyrrolidin-1-ylpropoxy, pyrrolidin-3-yloxy, pyrrolidin-2-ylmethoxy, 2-pyrrolidin-2-ylethoxy, 3-pyrrolidin-2-ylpropoxy, 2-hydroxy-3-morpholinopropoxy, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)ethoxy, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-hydroxy-3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 20 2-hydroxy-3-piperidinopropoxy, piperidin-3-yloxy, piperidin-4-yloxy, piperidin-3-ylmethoxy, 2-piperidin-3-ylethoxy, piperidin-4-ylmethoxy, 2-piperidin-4-ylethoxy, 3-homopiperidin-1-ylpropoxy, 3-homopiperidin-1-yl-2-hydroxypropoxy, 2-hydroxy-3-piperazin-1-ylpropoxy, 3-homopiperazin-1-ylpropoxy or 2-hydroxy-3-homopiperazin-1-ylpropoxy,

25 or R<sup>2</sup> is a group selected from 2-hydroxy-3-methylaminopropoxy, 3-ethylamino-2-hydroxypropoxy, 2-hydroxy-3-isopropylaminopropoxy, 3-dimethylamino-2-hydroxypropoxy, 3-diethylamino-2-hydroxypropoxy, 3-(N-ethyl-N-isopropylamino)-2-hydroxypropoxy, 3-(N-ethyl-N-methylamino)-2-hydroxypropoxy or 3-(N-isopropyl-N-methylamino)-2-hydroxypropoxy,

30 and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a R<sup>2</sup> substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, cyano, amino, vinyl, ethynyl, methoxy, methylsulphonyl, methylamino and dimethylamino,

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and wherein any heteroaryl or heterocyclyl group within a substituent on  $R^2$  optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, cyano, hydroxy, amino, methyl, ethyl, cyclopropyl, allyl, methoxy and acetyl,

and wherein any heterocyclyl group within a substituent on  $R^2$  optionally bears 1 or 2  
5 oxo substituents; and

$R^3$  is chloro or bromo;

or a pharmaceutically-acceptable acid-addition salt thereof.

A further particular compound of the invention is a quinazoline derivative of the Formula I wherein :

10  $R^1$  is methoxy and  $R^2$  is 4-pyridylmethoxy, 2-pyrid-4-yloxyethoxy, 3-pyrid-2-yloxypropoxy, 3-pyrid-4-yloxypropoxy, 3-azetidin-1-yl-2-hydroxypropoxy, 2-pyrrolidin-1-ylethoxy, 3-pyrrolidin-1-ylpropoxy, 2-hydroxy-3-pyrrolidin-1-ylpropoxy, 2-hydroxy-3-morpholinopropoxy, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-hydroxy-3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-hydroxy-  
15 3-piperidinopropoxy, piperidin-4-ylmethoxy, 2-piperidin-4-ylethoxy, 3-homopiperidin-1-yl-2-hydroxypropoxy or 2-hydroxy-3-piperazin-1-ylpropoxy,

or  $R^2$  is a group selected from 3-dimethylamino-2-hydroxypropoxy, 3-diethylamino-2-hydroxypropoxy, 3-(N-ethyl-N-isopropylamino)-2-hydroxypropoxy, 3-(N-ethyl-N-methylamino)-2-hydroxypropoxy or 3-(N-isopropyl-N-methylamino)-2-hydroxypropoxy,

20 and wherein any  $CH_2$  or  $CH_3$  group within a  $R^2$  substituent optionally bears on each said  $CH_2$  or  $CH_3$  group a substituent selected from hydroxy, cyano, amino, vinyl, ethynyl, methoxy, methylsulphonyl, methylamino and dimethylamino,

and wherein any pyridyl or heterocyclyl group within a substituent on  $R^2$  optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, cyano, hydroxy, amino, methyl, ethyl, cyclopropyl, allyl, methoxy and acetyl,  
25 and wherein any heterocyclyl group within a substituent on  $R^2$  optionally bears 1 or 2 oxo substituents; and

$R^3$  is chloro or bromo;

or a pharmaceutically-acceptable acid-addition salt thereof.

30 A particular compound of the invention is, for example, a quinazoline derivative of the Formula I selected from :-

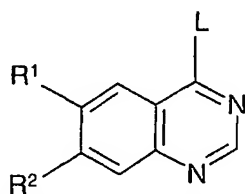
4-(2-chloro-5-methoxyanilino)-6-methoxy-7-(N-methylpiperidin-4-ylmethoxy)quinazoline,

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4-(2-chloro-5-methoxyanilino)-6-methoxy-7-piperidin-4-ylmethoxyquinazoline and  
 4-(2-bromo-5-methoxyanilino)-6-methoxy-7-[2-(N-methylpiperidin-4-yl)ethoxy]quinazoline;  
 or a pharmaceutically-acceptable acid-addition salt thereof.

A quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt  
 thereof, may be prepared by any process known to be applicable to the preparation of  
 chemically-related compounds. Such processes, when used to prepare a quinazoline  
 derivative of the Formula I are provided as a further feature of the invention and are illustrated  
 by the following representative process variants in which, unless otherwise stated, R<sup>1</sup>, R<sup>2</sup> and  
 R<sup>3</sup> have any of the meanings defined hereinbefore. Necessary starting materials may be  
 obtained by standard procedures of organic chemistry. The preparation of such starting  
 materials is described in conjunction with the following representative process variants and  
 within the accompanying Examples. Alternatively necessary starting materials are obtainable  
 by analogous procedures to those illustrated which are within the ordinary skill of an organic  
 chemist.

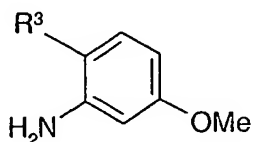
- (a) The reaction, conveniently in the presence of a suitable acid or base, of a quinazoline  
 of the Formula II



II

wherein L is a displaceable group and R<sup>1</sup> and R<sup>2</sup> have any of the meanings defined  
 hereinbefore except that any functional group is protected if necessary, with an aniline of the

Formula III



III

wherein R<sup>3</sup> has any of the meanings defined hereinbefore except that any functional group is  
 protected if necessary, whereafter any protecting group that is present is removed by  
 conventional means.

- A suitable displaceable group L is, for example, a halogeno, alkoxy, aryloxy or  
 sulphonyloxy group, for example a chloro, bromo, methoxy, phenoxy, pentafluorophenoxy,  
 methanesulphonyloxy or toluene-4-sulphonyloxy group.

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A suitable base is, for example, an organic amine base such as, for example, pyridine, 2,6-lutidine, collidine, 4-dimethylaminopyridine, triethylamine, morpholine, N-methyldmorpholine or diazabicyclo[5.4.0]undec-7-ene, or, for example, an alkali or alkaline earth metal carbonate or hydroxide, for example sodium carbonate, potassium carbonate, calcium carbonate, sodium hydroxide or potassium hydroxide, or, for example, an alkali metal hydride, for example sodium hydride.

The reaction is conveniently carried out in the presence of a suitable inert solvent or diluent, for example an alcohol or ester such as methanol, ethanol, isopropanol or ethyl acetate, a halogenated solvent such as methylene chloride, chloroform or carbon tetrachloride, an ether such as tetrahydrofuran or 1,4-dioxan, an aromatic solvent such as toluene, or a dipolar aprotic solvent such as N,N-dimethylformamide, N,N-dimethylacetamide, N-methylpyrrolidin-2-one or dimethylsulphoxide. The reaction is conveniently carried out at a temperature in the range, for example, 10 to 250°C, preferably in the range 40 to 80°C.

Typically, the quinazoline of the Formula II may be reacted with an aniline of the Formula III in the presence of a protic solvent such as isopropanol, conveniently in the presence of a suitable acid, for example hydrogen chloride gas in diethyl ether, or hydrochloric acid, and at a temperature in the range, for example, 0 to 150°C, preferably at or near the reflux temperature of the reaction solvent.

The quinazoline derivative of the Formula I may be obtained from this process in the form of the free base or alternatively it may be obtained in the form of a salt with the acid of the formula H-L wherein L has the meaning defined hereinbefore. When it is desired to obtain the free base from the salt, the salt may be treated with a suitable base, for example, an organic amine base such as, for example, pyridine, 2,6-lutidine, collidine, 4-dimethylaminopyridine, triethylamine, morpholine, N-methyldmorpholine or diazabicyclo[5.4.0]undec-7-ene, or, for example, an alkali or alkaline earth metal carbonate or hydroxide, for example sodium carbonate, potassium carbonate, calcium carbonate, sodium hydroxide or potassium hydroxide.

Protecting groups may in general be chosen from any of the groups described in the literature or known to the skilled chemist as appropriate for the protection of the group in question and may be introduced by conventional methods. Protecting groups may be removed by any convenient method as described in the literature or known to the skilled chemist as appropriate for the removal of the protecting group in question, such methods being chosen so

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as to effect removal of the protecting group with minimum disturbance of groups elsewhere in the molecule.

Specific examples of protecting groups are given below for the sake of convenience, in which "lower", as in, for example, lower alkyl, signifies that the group to which it is applied preferably has 1-4 carbon atoms. It will be understood that these examples are not exhaustive. Where specific examples of methods for the removal of protecting groups are given below these are similarly not exhaustive. The use of protecting groups and methods of deprotection not specifically mentioned are, of course, within the scope of the invention.

A carboxy protecting group may be the residue of an ester-forming aliphatic or arylaliphatic alcohol or of an ester-forming silanol (the said alcohol or silanol preferably containing 1-20 carbon atoms). Examples of carboxy protecting groups include straight or branched chain (1-12C)alkyl groups (for example isopropyl, and tert-butyl); lower alkoxy- lower alkyl groups (for example methoxymethyl, ethoxymethyl and isobutoxymethyl); lower acyloxy-lower alkyl groups, (for example acetoxymethyl, propionyloxymethyl, butyryloxymethyl and pivaloyloxymethyl); lower alkoxy-carbonyloxy-lower alkyl groups (for example 1-methoxycarbonyloxyethyl and 1-ethoxycarbonyloxyethyl); aryl-lower alkyl groups (for example benzyl, 4-methoxybenzyl, 2-nitrobenzyl, 4-nitrobenzyl, benzhydryl and phthalidyl); tri(lower alkyl)silyl groups (for example trimethylsilyl and tert-butyldimethylsilyl); tri(lower alkyl)silyl-lower alkyl groups (for example trimethylsilylethyl); and (2-6C)alkenyl groups (for example allyl). Methods particularly appropriate for the removal of carboxyl protecting groups include for example acid-, base-, metal- or enzymically-catalysed cleavage.

Examples of hydroxy protecting groups include lower alkyl groups (for example tert-butyl), lower alkenyl groups (for example allyl); lower alkanoyl groups (for example acetyl); lower alkoxy-carbonyl groups (for example tert-butoxycarbonyl); lower alkenyloxycarbonyl groups (for example allyloxycarbonyl); aryl-lower alkoxy-carbonyl groups (for example benzyloxycarbonyl, 4-methoxybenzyloxycarbonyl, 2-nitrobenzyloxycarbonyl and 4-nitrobenzyloxycarbonyl); tri(lower alkyl)silyl (for example trimethylsilyl and tert-butyldimethylsilyl) and aryl-lower alkyl (for example benzyl) groups.

Examples of amino protecting groups include formyl, aryl-lower alkyl groups (for example benzyl and substituted benzyl, 4-methoxybenzyl, 2-nitrobenzyl and 2,4-dimethoxybenzyl, and triphenylmethyl); di-4-anisylmethyl and furylmethyl groups; lower alkoxy-carbonyl (for example tert-butoxycarbonyl); lower alkenyloxycarbonyl (for example

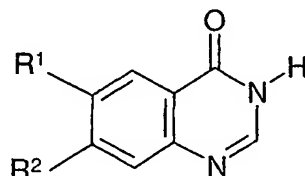
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allyloxycarbonyl); aryl-lower alkoxy carbonyl groups (for example benzyloxycarbonyl, 4-methoxybenzyloxycarbonyl, 2-nitrobenzyloxycarbonyl and 4-nitrobenzyloxycarbonyl); trialkylsilyl (for example trimethylsilyl and tert-butyldimethylsilyl); alkylidene (for example methyldiene) and benzylidene and substituted benzylidene groups.

5 Methods appropriate for removal of hydroxy and amino protecting groups include, for example, acid-, base-, metal- or enzymically-catalysed hydrolysis for groups such as 2-nitrobenzyloxycarbonyl, hydrogenation for groups such as benzyl and photolytically for groups such as 2-nitrobenzyloxycarbonyl.

The reader is referred to Advanced Organic Chemistry, 4th Edition, by J. March, 10 published by John Wiley & Sons 1992, for general guidance on reaction conditions and reagents and to Protective Groups in Organic Synthesis, 2<sup>nd</sup> Edition, by T. Green *et al.*, also published by John Wiley & Son, for general guidance on protecting groups.

Quinazoline starting materials of the Formula II may be obtained by conventional procedures. For example, a 3,4-dihydroquinazolin-4-one of Formula IV

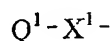


IV

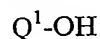
15 wherein R<sup>1</sup> and R<sup>2</sup> have any of the meanings defined hereinbefore except that any functional group is protected if necessary, may be reacted with a halogenating agent such as thionyl chloride, phosphoryl chloride or a mixture of carbon tetrachloride and triphenylphosphine whereafter any protecting group that is present is removed by conventional means.

20 The 4-chloroquinazoline so obtained may be converted, if required, into a 4-pentafluorophenoxyquinazoline by reaction with pentafluorophenol in the presence of a suitable base such as potassium carbonate and in the presence of a suitable solvent such as N,N-dimethylformamide.

(b) For the production of those compounds of the Formula I wherein R<sup>2</sup> is a group of the 25 formula :

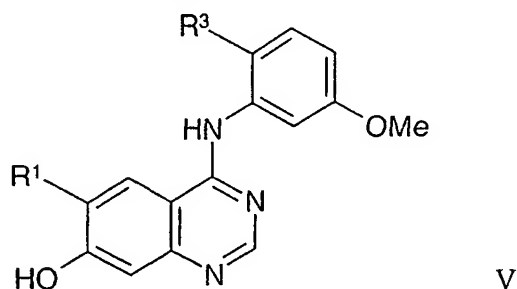


wherein X<sup>1</sup> is an oxygen atom, the coupling, conveniently in the presence of a suitable dehydrating agent, of an alcohol of the Formula



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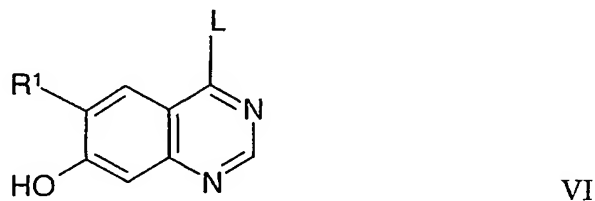
wherein  $Q^1$  has any of the meanings defined hereinbefore except that any functional group is protected if necessary, with a quinazoline of the Formula V



wherein  $R^1$  and  $R^3$  have any of the meanings defined hereinbefore except that any functional group is protected if necessary, whereafter any protecting group that is present is removed by conventional means.

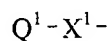
A suitable dehydrating agent is, for example, a carbodiimide reagent such as dicyclohexylcarbodiimide or 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide or a mixture of an azo compound such as diethyl or di-tert-butyl azodicarboxylate and a phosphine such as triphenylphosphine. The reaction is conveniently carried out in the presence of a suitable inert solvent or diluent, for example a halogenated solvent such as methylene chloride, chloroform or carbon tetrachloride and at a temperature in the range, for example, 10 to 150°C, preferably at or near ambient temperature.

The quinazoline of the Formula V may be obtained by conventional procedures. For example, a quinazoline of the Formula VI



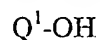
wherein L is a displaceable group as defined hereinbefore and  $R^1$  has any of the meanings defined hereinbefore except that any functional group is protected if necessary, may be reacted with an aniline of the Formula III as defined hereinbefore except that any functional group is protected if necessary, whereafter any protecting group that is present is removed by conventional means.

(c) For the production of those compounds of the Formula I wherein  $R^1$  is a group of the formula :

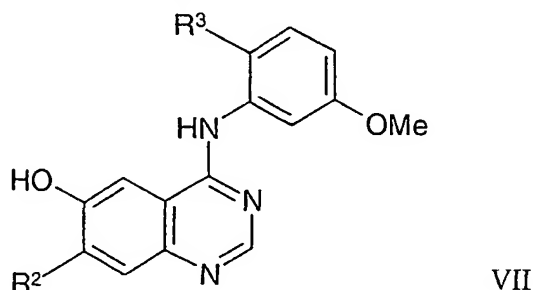


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wherein  $X^1$  is an oxygen atom, the coupling, conveniently in the presence of a suitable dehydrating agent as defined hereinbefore, of an alcohol of the Formula



wherein  $Q^1$  has any of the meanings defined hereinbefore except that any functional group is  
 5 protected if necessary, with a quinazoline of the Formula VII



wherein  $R^2$  and  $R^3$  have any of the meanings defined hereinbefore except that any functional group is protected if necessary, whereafter any protecting group that is present is removed by conventional means.

10 The reaction is conveniently carried out in the presence of a suitable inert solvent or diluent, for example a halogenated solvent such as methylene chloride, chloroform or carbon tetrachloride and at a temperature in the range, for example, 10 to 150°C, preferably at or near ambient temperature.

The quinazoline of the Formula VII may be obtained by conventional procedures  
 15 analogous to those described hereinbefore for the preparation of the quinazoline of the Formula V.

(d) For the production of those compounds of the Formula I wherein  $R^1$  or  $R^2$  contains an amino-hydroxy-disubstituted (1-6C)alkoxy group (such as 2-hydroxy-3-piperidinopropoxy, 2-hydroxy-3-methylaminopropoxy, 3-dimethylamino-2-hydroxypropoxy or  
 20 3-[N-(3-dimethylaminopropyl)-N-methylamino]-2-hydroxypropoxy), the reaction of a compound of the Formula I wherein  $R^1$  or  $R^2$  contains an epoxy-substituted (1-6C)alkoxy group with a heterocyclyl compound or an appropriate amine.

The reaction is conveniently carried out in the presence of a suitable inert diluent or carrier as defined hereinbefore and at a temperature in the range 10 to 150°C, preferably at or  
 25 near ambient temperature.

(e) For the production of those compounds of the Formula I wherein  $R^1$  or  $R^2$  contains an amino-acyloxy-disubstituted (1-6C)alkoxy group (such as a 2-isobutyryloxy-3-pyrrolidin-



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1-ylpropoxy group), the acylation of a compound of the Formula I wherein R<sup>1</sup> or R<sup>2</sup> contains an amino-hydroxy-disubstituted (1-6C)alkoxy group.

The acylation reaction is conveniently carried out in the presence of a suitable inert diluent or carrier as defined hereinbefore and at a temperature in the range 10 to 150°C, preferably at or near ambient temperature. For example, the acylation reaction may be carried out by the reaction of a compound of the Formula I wherein R<sup>1</sup> or R<sup>2</sup> contains an amino-hydroxy-disubstituted (1-6C)alkoxy group with an appropriate carboxylic acid, conveniently in the presence of a suitable dehydrating agent as defined hereinbefore.

(f) For the production of those compounds of the Formula I wherein an R<sup>1</sup> or R<sup>2</sup> group contains a hydroxy group, the cleavage of the corresponding compound of the Formula I wherein the R<sup>1</sup> or R<sup>2</sup> group contains a protected hydroxy group.

Suitable protecting groups for a hydroxy group are, for example, any of the protecting groups disclosed hereinbefore. Suitable methods for the cleavage of such hydroxy protecting groups are also disclosed hereinbefore. In particular, a suitable protecting group is a lower alkanoyl group such as an acetyl group which may be cleaved under conventional reaction conditions such as under base-catalysed conditions, for example in the presence of ammonia.

When a pharmaceutically-acceptable salt of a quinazoline derivative of the Formula I is required, for example an acid-addition salt, it may be obtained by, for example, reaction of said quinazoline derivative with a suitable acid using a conventional procedure.

## 20 Biological Assays

The following assays can be used to measure the effects of the compounds of the present invention as c-Src tyrosine kinase inhibitors, as inhibitors in vitro of the proliferation of c-Src transfected fibroblast cells, as inhibitors in vitro of the migration of A549 human lung tumour cells and as inhibitors in vivo of the growth in nude mice of xenografts of A549 tissue.

### 25 (a) In Vitro Enzyme Assay

The ability of test compounds to inhibit the phosphorylation of a tyrosine containing polypeptide substrate by the enzyme c-Src kinase was assessed using a conventional Elisa assay.

A substrate solution [100µl of a 20µg/ml solution of the polyamino acid Poly(Glu, Tyr) 4:1 (Sigma Catalogue No. P0275) in phosphate buffered saline (PBS) containing 0.2mg/ml of sodium azide] was added to each well of a number of Nunc 96-well immunoplates (Catalogue No. 439454) and the plates were sealed and stored at 4°C for

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16 hours. The excess of substrate solution was discarded, and aliquots of Bovine Serum Albumin (BSA; 150µl of a 5% solution in PBS) were transferred into each substrate-coated assay well and incubated for 1 hour at ambient temperature to block non specific binding. The assay plate wells were washed in turn with PBS containing 0.05% v/v Tween 20 (PBST) and  
5 with Hepes pH7.4 buffer (50mM, 300µl/well) before being blotted dry.

Each test compound was dissolved in dimethyl sulphoxide and diluted with distilled water to give a series of dilutions (from 100µM to 0.001µM). Portions (25µl) of each dilution of test compound were transferred to wells in the washed assay plates. "Total" control wells contained diluted DMSO instead of compound. Aliquots (25µl) of an aqueous magnesium  
10 chloride solution (80mM) containing adenosine-5'-triphosphate (ATP; 40µM) was added to all test wells except the "blank" control wells which contained magnesium chloride without ATP.

Active human c-Src kinase (recombinant enzyme expressed in Sf9 insect cells; obtained from Upstate Biotechnology Inc. product 14-117) was diluted immediately prior to  
15 use by a factor of 1:10,000 with an enzyme diluent which comprised 100mM Hepes pH7.4 buffer, 0.2mM sodium orthovanadate, 2mM dithiothreitol and 0.02% BSA. To start the reactions, aliquots (50µl) of freshly diluted enzyme were added to each well and the plates were incubated at ambient temperature for 20 minutes. The supernatant liquid in each well was discarded and the wells were washed twice with PBST. Mouse IgG anti-phosphotyrosine  
20 antibody (Upstate Biotechnology Inc. product 05-321; 100µl) was diluted by a factor of 1:6000 with PBST containing 0.5% w/v BSA and added to each well. The plates were incubated for 1 hour at ambient temperature. The supernatant liquid was discarded and each well was washed with PBST (x4). Horse radish peroxidase (HRP)-linked sheep anti-mouse Ig antibody (Amersham Catalogue No. NXA 931; 100µl) was diluted by a factor of 1:500 with  
25 PBST containing 0.5% w/v BSA and added to each well. The plates were incubated for 1 hour at ambient temperature. The supernatant liquid was discarded and the wells were washed with PBST (x4).

A PCSB capsule (Sigma Catalogue No. P4922) was dissolved in distilled water (100ml) to provide phosphate-citrate pH5 buffer (50mM) containing 0.03% sodium perborate.  
30 An aliquot (50ml) of this buffer was mixed with a 50mg tablet of 2,2'-azinobis(3-ethylbenzothiazoline-6-sulphonic acid) (ABTS; Boehringer Catalogue No. 1204 521). Aliquots (100µl) of the resultant solution were added to each well. The plates

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were incubated for 20 to 60 minutes at ambient temperature until the optical density value of the "total" control wells, measured at 405nm using a plate reading spectrophotometer, was approximately 1.0. "Blank" (no ATP) and "total" (no compound) control values were used to determine the dilution range of test compound which gave 50% inhibition of enzyme activity.

5 (b) In Vitro c-Src transfected NIH 3T3 (c-src 3T3) Fibroblast Proliferation Assay

This assay determined the ability of a test compound to inhibit the proliferation of National Institute of Health (NIH) mouse 3T3 fibroblast cells that had been stably-transfected with an activating mutant (Y530F) of human c-Src.

Using a similar procedure to that described by Shalloway *et al.*, Cell, 1987, 49, 65-73,  
10 NIH 3T3 cells were transfected with an activating mutant (Y530F) of human c-Src. The resultant c-Src 3T3 cells were typically seeded at  $1.5 \times 10^4$  cells per well into 96-well tissue-culture-treated clear assay plates (Costar) each containing an assay medium comprising Dulbecco's modified Eagle's medium (DMEM; Sigma) plus 0.5% foetal calf serum (FCS), 2mM glutamine, 100 units/ml penicillin and 0.1mg/ml streptomycin in 0.9% aqueous sodium  
15 chloride solution. The plates were incubated overnight at 37°C in a humidified (7.5% CO<sub>2</sub> : 95% air) incubator.

Test compounds were solubilised in DMSO to form a 10mM stock solution. Aliquots of the stock solution were diluted with the DMEM medium described above and added to appropriate wells. Serial dilutions were made to give a range of test concentrations. Control  
20 wells to which test compound was not added were included on each plate. The plates were incubated overnight at 37°C in a humidified (7.5% CO<sub>2</sub> : 95% air) incubator.

BrdU labelling reagent (Boehringer Mannheim Catalogue No. 647 229) was diluted by a factor of 1:100 in DMEM medium containing 0.5% FCS and aliquots (20µl) were added to each well to give a final concentration of 10µM. The plates were incubated at 37°C for  
25 2 hours. The medium was decanted. A denaturing solution (FixDenat solution, Boehringer Mannheim Catalogue No. 647 229; 50µl) was added to each well and the plates were placed on a plate shaker at ambient temperature for 45 minutes. The supernatant was decanted and the wells were washed with PBS (200µl per well). Anti-BrdU-Peroxidase solution (Boehringer Mannheim Catalogue No. 647 229) was diluted by a factor of 1:100 in PBS  
30 containing 1% BSA and 0.025% dried skimmed milk (Marvel (registered trade mark), Premier Beverages, Stafford, GB) and an aliquot (100µl) of the resultant solution was added to each well. The plates were placed on a plate shaker at ambient temperature for 90 minutes. The

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wells were washed with PBS (x5) to ensure removal of non bound antibody conjugate. The plates were blotted dry and tetramethylbenzidine substrate solution (Boehringer Mannheim Catalogue No. 647 229; 100µl) was added to each well. The plates were gently agitated on a plate shaker while the colour developed during a 10 to 20 minute period. The absorbance of the wells was measured at 690nm. The extent of inhibition of cellular proliferation at a range of concentrations of each test compound was determined and an anti-proliferative IC<sub>50</sub> value was derived.

(c) In Vitro Microdroplet Migration Assay

This assay determines the ability of a test compound to inhibit the migration of adherent mammalian cell lines, for example the human tumour cell line A549.

RPMI medium(Sigma) containing 10% FCS, 1% L-glutamine and 0.3% agarose (Difco Catalogue No. 0142-01) was warmed to 37°C in a waterbath. A stock 2% aqueous agar solution was autoclaved and stored at 42°C. An aliquot (1.5 ml) of the agar solution was added to RPMI medium (10 ml) immediately prior to its use. A549 cells (Accession No. ATCC CCL185) were suspended at a concentration of  $2 \times 10^7$  cells/ml in the medium and maintained at a temperature of 37°C.

A droplet (2µl) of the cell/agarose mixture was transferred by pipette into the centre of each well of a number of 96-well, flat bottomed non-tissue-culture-treated microtitre plate (Bibby Sterilin Catalogue No. 642000). The plates were placed briefly on ice to speed the gelling of the agarose-containing droplets. Aliquots (90µl) of medium which had been cooled to 4°C were transferred into each well, taking care not to disturb the microdroplets. Test compounds were diluted from a 10mM stock solution in DMSO using RPMI medium as described above. Aliquots (10µl) of the diluted test compounds were transferred to the wells, again taking care not to disturb the microdroplets. The plates were incubated at 37°C in a humidified (7.5% CO<sub>2</sub> : 95% air) incubator for about 48 hours.

Migration was assessed visually and the distance of migration was measured back to the edge of the agar droplet. A migratory inhibitory IC<sub>50</sub> was derived by plotting the mean migration measurement against test compound concentration.

(d) In Vivo A549 Xenograft Growth Assay

This test measures the ability of compounds to inhibit the growth of the A549 human carcinoma grown as a tumour in athymic nude mice (Alderley Park nu/nu strain). A total of about  $5 \times 10^6$  A549 cells in matrigel (Beckton Dickinson Catalogue No. 40234) were injected

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subcutaneously into the left flank of each test mouse and the resultant tumours were allowed to grow for about 14 days. Tumour size was measured twice weekly using callipers and a theoretical volume was calculated. Animals were selected to provide control and treatment groups of approximately equal average tumour volume. Test compounds were prepared as a ball-milled suspension in 1% polysorbate vehicle and dosed orally once daily for a period of about 28 days. The effect on tumour growth was assessed.

Although the pharmacological properties of the compounds of the Formula I vary with structural change as expected, in general activity possessed by compounds of the Formula I, may be demonstrated at the following concentrations or doses in one or more of the above tests (a), (b), (c) and (d):-

Test (a):-  $IC_{50}$  in the range, for example, 0.001 - 10  $\mu$ M;

Test (b):-  $IC_{50}$  in the range, for example, 0.01 - 20  $\mu$ M;

Test (c):- activity in the range, for example, 0.01-25  $\mu$ M;

Test (d):- activity in the range, for example, 1-200 mg/kg/day.

No physiologically-unacceptable toxicity was observed in Test (d) at the effective dose for compounds tested of the present invention. Accordingly no untoward toxicological effects are expected when a compound of Formula I, or a pharmaceutically-acceptable salt thereof, as defined hereinbefore is administered at the dosage ranges defined hereinafter.

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, as defined hereinbefore in association with a pharmaceutically-acceptable diluent or carrier.

The compositions of the invention may be in a form suitable for oral use (for example as tablets, lozenges, hard or soft capsules, aqueous or oily suspensions, emulsions, dispersible powders or granules, syrups or elixirs), for topical use (for example as creams, ointments, gels, or aqueous or oily solutions or suspensions), for administration by inhalation (for example as a finely divided powder or a liquid aerosol), for administration by insufflation (for example as a finely divided powder) or for parenteral administration (for example as a sterile aqueous or oily solution for intravenous, subcutaneous, intramuscular or intramuscular dosing or as a suppository for rectal dosing).

The compositions of the invention may be obtained by conventional procedures using conventional pharmaceutical excipients, well known in the art. Thus, compositions intended for oral use may contain, for example, one or more colouring, sweetening, flavouring and/or

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preservative agents.

The amount of active ingredient that is combined with one or more excipients to produce a single dosage form will necessarily vary depending upon the host treated and the particular route of administration. For example, a formulation intended for oral  
5 administration to humans will generally contain, for example, from 0.5 mg to 0.5 g of active agent (more suitably from 0.5 to 100 mg, for example from 1 to 30 mg) compounded with an appropriate and convenient amount of excipients which may vary from about 5 to about 98 percent by weight of the total composition.

The size of the dose for therapeutic or prophylactic purposes of a compound of the  
10 Formula I will naturally vary according to the nature and severity of the conditions, the age and sex of the animal or patient and the route of administration, according to well known principles of medicine.

In using a compound of the Formula I for therapeutic or prophylactic purposes it will generally be administered so that a daily dose in the range, for example, 0.1 mg/kg to  
15 75 mg/kg body weight is received, given if required in divided doses. In general lower doses will be administered when a parenteral route is employed. Thus, for example, for intravenous administration, a dose in the range, for example, 0.1 mg/kg to 30 mg/kg body weight will generally be used. Similarly, for administration by inhalation, a dose in the range, for example, 0.05 mg/kg to 25 mg/kg body weight will be used. Oral administration is however  
20 preferred, particularly in tablet form. Typically, unit dosage forms will contain about 0.5 mg to 0.5 g of a compound of this invention.

According to a further aspect of the invention there is provided a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, as defined hereinbefore for use in a method of treatment of the human or animal body by therapy.

25 As stated above, it is known that the predominant role of c-Src non-receptor tyrosine kinase is to regulate cell motility which is necessarily required for a localised tumour to progress through the stages of dissemination into the blood stream, invasion of other tissues and initiation of metastatic tumour growth. We have found that the quinazoline derivatives of the present invention possess potent anti-tumour activity which it is believed is obtained by  
30 way of inhibition of one or more of the non-receptor tyrosine-specific protein kinases such as c-Src kinase that are involved in the signal transduction steps which lead to the invasiveness and migratory ability of metastasising tumour cells.

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Accordingly the quinazoline derivatives of the present invention are of value as anti-tumour agents, in particular as selective inhibitors of the motility, dissemination and invasiveness of mammalian cancer cells leading to inhibition of metastatic tumour growth. Particularly, the quinazoline derivatives of the present invention are of value as anti-invasive agents in the containment and/or treatment of solid tumour disease. Particularly, the compounds of the present invention are expected to be useful in the prevention or treatment of those tumours which are sensitive to inhibition of one or more of the multiple non-receptor tyrosine kinases such as c-Src kinase that are involved in the signal transduction steps which lead to the invasiveness and migratory ability of metastasising tumour cells. Further, the compounds of the present invention are expected to be useful in the prevention or treatment of those tumours which are mediated alone or in part by inhibition of the enzyme c-Src, *i.e.* the compounds may be used to produce a c-Src enzyme inhibitory effect in a warm-blooded animal in need of such treatment. Specifically, the compounds of the present invention are expected to be useful in the prevention or treatment of solid tumour disease.

Thus according to this aspect of the invention there is provided the use of a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, as defined hereinbefore in the manufacture of a medicament for use as an anti-invasive agent in the containment and/or treatment of solid tumour disease.

According to a further feature of this aspect of the invention there is provided a method for producing an anti-invasive effect by the containment and/or treatment of solid tumour disease in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, as defined hereinbefore.

According to a further aspect of the invention there is provided the use of a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, as defined hereinbefore in the manufacture of a medicament for use in the prevention or treatment of solid tumour disease in a warm-blooded animal such as man.

According to a further feature of this aspect of the invention there is provided a method for the prevention or treatment of solid tumour disease in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, as defined hereinbefore.

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According to a further aspect of the invention there is provided the use of a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, as defined hereinbefore in the manufacture of a medicament for use in the prevention or treatment of those tumours which are sensitive to inhibition of non-receptor tyrosine kinases  
5 such as c-Src kinase that are involved in the signal transduction steps which lead to the invasiveness and migratory ability of metastasising tumour cells.

According to a further feature of this aspect of the invention there is provided a method for the prevention or treatment of those tumours which are sensitive to inhibition of non-receptor tyrosine kinases such as c-Src kinase that are involved in the signal transduction  
10 steps which lead to the invasiveness and migratory ability of metastasising tumour cells which comprises administering to said animal an effective amount of a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, as defined hereinbefore.

According to a further aspect of the invention there is provided the use of a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, as  
15 defined hereinbefore in the manufacture of a medicament for use in providing a c-Src kinase inhibitory effect.

According to a further feature of this aspect of the invention there is provided a method for providing a c-Src kinase inhibitory effect which comprises administering to said animal an effective amount of a quinazoline derivative of the Formula I, or a  
20 pharmaceutically-acceptable salt thereof, as defined hereinbefore.

The anti-invasive treatment defined hereinbefore may be applied as a sole therapy or may involve, in addition to the quinazoline derivative of the invention, conventional surgery or radiotherapy or chemotherapy. Such chemotherapy may include one or more of the following categories of anti-tumour agents :-

- 25 (i) other anti-invasion agents (for example metalloproteinase inhibitors like marimastat and inhibitors of urokinase plasminogen activator receptor function);  
(ii) antiproliferative/antineoplastic drugs and combinations thereof, as used in medical oncology, such as alkylating agents (for example cis-platin, carboplatin, cyclophosphamide, nitrogen mustard, melphalan, chlorambucil, busulphan and nitrosoureas); antimetabolites (for  
30 example antifolates such as fluoropyrimidines like 5-fluorouracil and tegafur, raltitrexed, methotrexate, cytosine arabinoside and hydroxyurea, or, for example, one of the preferred antimetabolites disclosed in European Patent Application No. 562734 such as  
(2S)-2-{o-fluoro-p-[N-(2,7-dimethyl-4-oxo-3,4-dihydroquinazolin-6-yl)methyl]-



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- N-(prop-2-ynyl)amino]benzamido}-4-(tetrazol-5-yl)butyric acid); antitumour antibiotics (for example anthracyclines like adriamycin, bleomycin, doxorubicin, daunomycin, epirubicin, idarubicin, mitomycin-C, dactinomycin and mithramycin); antimitotic agents (for example vinca alkaloids like vincristine, vinblastine, vindesine and vinorelbine and taxoids like taxol and taxotere); and topoisomerase inhibitors (for example epipodophyllotoxins like etoposide and teniposide, amsacrine, topotecan and camptothecin);
- (iii) cytostatic agents such as antioestrogens (for example tamoxifen, toremifene, raloxifene, droloxifene and idoxifyfene), antiandrogens (for example bicalutamide, flutamide, nilutamide and cyproterone acetate), LHRH antagonists or LHRH agonists (for example goserelin, leuporelin and buserelin), progestogens (for example megestrol acetate), aromatase inhibitors (for example as anastrozole, letrozole, vorazole and exemestane) and inhibitors of 5 $\alpha$ -reductase such as finasteride;
- (iv) inhibitors of growth factor function, for example such inhibitors include growth factor antibodies, growth factor receptor antibodies, tyrosine kinase inhibitors and serine/threonine kinase inhibitors, for example inhibitors of the epidermal growth factor family (for example the EGFR tyrosine kinase inhibitors N-(3-chloro-4-fluorophenyl)-7-methoxy-6-(3-morpholinopropoxy)quinazolin-4-amine (ZD1839), N-(3-ethynylphenyl)-6,7-bis(2-methoxyethoxy)quinazolin-4-amine (CP 358774) and 6-acrylamido-N-(3-chloro-4-fluorophenyl)-7-(3-morpholinopropoxy)quinazolin-4-amine (CI 1033)), for example inhibitors of the platelet-derived growth factor family and for example inhibitors of the hepatocyte growth factor family; and
- (v) antiangiogenic agents such as those which inhibit vascular endothelial growth factor such as the compounds disclosed in International Patent Applications WO 97/22596, WO 97/30035, WO 97/32856 and WO 98/13354 and those that work by other mechanisms (for example linomide, inhibitors of integrin  $\alpha v \beta 3$  function and angiostatin).

Such conjoint treatment may be achieved by way of the simultaneous, sequential or separate dosing of the individual components of the treatment. Such combination products employ the compounds of this invention within the dosage range described hereinbefore and the other pharmaceutically-active agent within its approved dosage range.

- According to this aspect of the invention there is provided a pharmaceutical product comprising a quinazoline derivative of the formula I as defined hereinbefore and an additional anti-tumour agent as defined hereinbefore for the conjoint treatment of cancer.

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Although the compounds of the Formula I are primarily of value as therapeutic agents for use in warm-blooded animals (including man), they are also useful whenever it is required to inhibit the effects of c-Src. Thus, they are useful as pharmacological standards for use in the development of new biological tests and in the search for new pharmacological agents.

5 The invention will now be illustrated in the following Examples in which, generally :

(i) operations were carried out at ambient temperature, *i.e.* in the range 17 to 25°C and under an atmosphere of an inert gas such as argon unless otherwise stated;

(ii) evaporations were carried out by rotary evaporation *in vacuo* and work-up procedures were carried out after removal of residual solids by filtration;

10 (iii) column chromatography (by the flash procedure) and medium pressure liquid chromatography (MPLC) were performed on Merck Kieselgel silica (Art. 9385) or Merck Lichroprep RP-18 (Art. 9303) reversed-phase silica obtained from E. Merck, Darmstadt, Germany or high pressure liquid chromatography (HPLC) was performed on C18 reverse phase silica, for example on a Dynamax C-18 60Å preparative reversed-phase column;

15 (iv) yields, where present, are not necessarily the maximum attainable;

(v) in general, the end-products of the Formula I have satisfactory microanalyses and their structures were confirmed by nuclear magnetic resonance (NMR) and/or mass spectral techniques; fast-atom bombardment (FAB) mass spectral data were obtained using a Platform spectrometer and, where appropriate, either positive ion data or negative ion data were  
20 collected; NMR chemical shift values were measured on the delta scale [proton magnetic resonance spectra were determined using a Jeol JNM EX 400 spectrometer operating at a field strength of 400MHz, Varian Gemini 2000 spectrometer operating at a field strength of 300MHz or a Bruker AM300 spectrometer operating at a field strength of 300MHz]; the following abbreviations have been used: s, singlet; d, doublet; t, triplet; q, quartet; m,  
25 multiplet; br, broad;

(vi) intermediates were not generally fully characterised and purity was assessed by thin layer chromatographic, HPLC, infra-red (IR) and/or NMR analysis;

(vii) melting points are uncorrected and were determined using a Mettler SP62 automatic melting point apparatus or an oil-bath apparatus; melting points for the  
30 end-products of the Formula I were determined after crystallisation from a conventional organic solvent such as ethanol, methanol, acetone, ether or hexane, alone or in admixture;

(viii) the following abbreviations have been used:-

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|      |                               |
|------|-------------------------------|
| DMF  | <u>N,N</u> -dimethylformamide |
| DMSO | dimethylsulphoxide            |
| THF  | tetrahydrofuran               |

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**Example 1** 4-(2-chloro-5-methoxyanilino)-6-methoxy-7-(N-methylpiperidin-4-ylmethoxy)quinazoline

A mixture of 4-chloro-6-methoxy-7-(N-methylpiperidin-4-ylmethoxy)quinazoline (0.18 g), 2-chloro-5-methoxyaniline (0.188 g), a 6.2M solution of hydrogen chloride in isopropanol (0.15 ml) and isopropanol (4 ml) was stirred and heated to 80°C for 2.5 hours. The mixture was cooled to ambient temperature and evaporated. The residue was triturated under diethyl ether. The solid so obtained was isolated, washed in turn with isopropanol and diethyl ether and dried under vacuum. The material so obtained was dissolved in methylene chloride and a saturated methanolic ammonia solution (0.5 ml) was added and the mixture was stirred at ambient temperature for 10 minutes. The resultant mixture was filtered, the filtrate was evaporated and the residue was purified by column chromatography on silica using a 19:1 mixture of methylene chloride and a saturated methanolic ammonia solution as eluent. The material so obtained was triturated under diethyl ether and the solid so obtained was isolated, washed with diethyl ether and dried under vacuum. There was thus obtained the title compound (0.07 g); NMR Spectrum: (DMSO-d<sub>6</sub>) 1.3-1.4 (m, 2H), 1.78 (m, 3H), 1.9 (m, 2H), 2.15 (s, 3H), 2.8 (d, 2H), 3.8 (s, 3H), 3.95 (s, 3H), 4.0 (d, 2H), 6.9 (d, 1H), 7.15 (m, 2H), 7.48 (d, 1H), 7.8 (s, 1H), 8.3 (s, 1H), 9.5 (br s, 1H); Mass Spectrum: M+H<sup>+</sup> 443 and 445.

The 4-chloro-6-methoxy-7-(N-methylpiperidin-4-ylmethoxy)quinazoline used as a starting material was prepared as follows :-

A solution of di-tert-butyl dicarbonate (41.7 g) in ethyl acetate (75 ml) was added dropwise to a stirred solution of ethyl piperidine-4-carboxylate (30 g) in ethyl acetate (150 ml) which had been cooled to 0 to 5°C in an ice-bath. The resultant mixture was stirred at ambient temperature for 48 hours. The mixture was poured into water (300 ml). The organic layer was separated, washed in turn with water (200 ml), 0.1N aqueous hydrochloric acid solution (200 ml), a saturated aqueous sodium bicarbonate solution (200 ml) and brine (200 ml), dried over magnesium sulphate and evaporated. There was thus obtained ethyl N-tert-butoxycarbonylpiperidine-4-carboxylate (48 g); NMR Spectrum: (CDCl<sub>3</sub>) 1.25 (t, 3H), 1.45 (s, 9H), 1.55-1.7 (m, 2H), 1.8-2.0 (d, 2H), 2.35-2.5 (m, 1H), 2.7-2.95 (t, 2H), 3.9-4.1 (br s, 2H), 4.15 (q, 2H).

A solution of the material so obtained in THF (180 ml) was cooled at 0°C and lithium aluminium hydride (1M solution in THF; 133 ml) was added dropwise. The mixture was stirred at 0°C for 2 hours. Water (30 ml) and 2N aqueous sodium hydroxide solution (10 ml) were added in turn and the mixture was stirred for 15 minutes. The resultant mixture was

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filtered through diatomaceous earth and the solids were washed with ethyl acetate. The filtrate was washed in turn with water and with brine, dried over magnesium sulphate and evaporated. There was thus obtained N-tert-butoxycarbonyl-4-hydroxymethylpiperidine (36.3 g); NMR Spectrum: (CDCl<sub>3</sub>) 1.05-1.2 (m, 2H), 1.35-1.55 (m, 10H), 1.6-1.8 (m, 2H),  
5 2.6-2.8 (t, 2H), 3.4-3.6 (t, 2H), 4.0-4.2 (br s, 2H).

1,4-Diazabicyclo[2.2.2]octane (42.4 g) was added to a solution of N-tert-butoxycarbonyl-4-hydroxymethylpiperidine (52.5 g) in tert-butyl methyl ether (525 ml) and the mixture was stirred at ambient temperature for 15 minutes. The mixture was then cooled in an ice-bath to 5°C and a solution of 4-toluenesulphonyl chloride (62.8 g) in  
10 tert-butyl methyl ether (525 ml) was added dropwise over 2 hours while maintaining the reaction temperature at approximately 0°C. The resultant mixture was allowed to warm to ambient temperature and was stirred for 1 hour. Petroleum ether (b.p. 60-80°C, 1L) was added and the precipitate was removed by filtration. The filtrate was evaporated to give a solid residue which was dissolved in diethyl ether. The organic solution was washed in turn  
15 with 0.5N aqueous hydrochloric acid solution, water, a saturated aqueous sodium bicarbonate solution and brine, dried over magnesium sulphate and evaporated. There was thus obtained N-tert-butoxycarbonyl-4-(4-toluenesulphonyloxymethyl)piperidine (76.7 g); NMR Spectrum: (CDCl<sub>3</sub>) 1.0-1.2 (m, 2H), 1.45 (s, 9H), 1.65 (d, 2H), 1.75-1.9 (m, 2H), 2.45 (s, 3H), 2.55-2.75 (m, 2H), 3.85 (d, 1H), 4.0-4.2 (br s, 2H), 7.35 (d, 2H), 7.8 (d, 2H).

20 A portion (40 g) of the material so obtained was added to a suspension of ethyl 4-hydroxy-3-methoxybenzoate (19.6 g) and potassium carbonate (28 g) in DMF (200 ml) and the resultant mixture was stirred and heated to 95°C for 2.5 hours. The mixture was cooled to ambient temperature and partitioned between water and a mixture of ethyl acetate and diethyl ether. The organic layer was washed in turn with water and brine, dried over magnesium  
25 sulphate and evaporated. The resulting oil was crystallised from petroleum ether (b.p. 60-80°C) and the suspension was stored overnight at 5°C. The resultant solid was collected by filtration, washed with petroleum ether and dried under vacuum. There was thus obtained ethyl 4-(N-tert-butoxycarbonylpiperidin-4-ylmethoxy)-3-methoxybenzoate (35 g); m.p. 81-83°C; NMR Spectrum: (CDCl<sub>3</sub>) 1.2-1.35 (m, 2H), 1.4 (t, 3H), 1.48 (s, 9H), 1.8-1.9 (d,  
30 2H), 2.0-2.15 (m, 2H), 2.75 (t, 2H), 3.9 (d, 2H), 3.95 (s, 3H), 4.05-4.25 (br s, 2H), 4.35 (q, 2H), 6.85 (d, 1H), 7.55 (s, 1H), 7.65 (d, 1H).

The material so obtained was dissolved in formic acid (35 ml), formaldehyde (12M, 37% in water, 35 ml) was added and the mixture was stirred and heated to 95°C for 3 hours.

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The resultant mixture was evaporated. The residue was dissolved in methylene chloride and hydrogen chloride (3M solution in diethyl ether; 40 ml) was added. The mixture was diluted with diethyl ether and the mixture was triturated until a solid was formed. The solid was collected, washed with diethyl ether and dried under vacuum overnight at 50°C. There was  
5 thus obtained ethyl 3-methoxy-4-(N-methylpiperidin-4-ylmethoxy)benzoate (30.6 g);  
NMR Spectrum: (DMSO-d<sub>6</sub>) 1.29 (t, 3H), 1.5-1.7 (m, 2H), 1.95 (d, 2H), 2.0-2.15 (br s, 1H), 2.72 (s, 3H), 2.9-3.1 (m, 2H), 3.35-3.5 (br s, 2H), 3.85 (s, 3H), 3.9-4.05 (br s, 2H), 4.3 (q, 2H), 7.1 (d, 1H), 7.48 (s, 1H), 7.6 (d, 1H).

The material so obtained was dissolved in methylene chloride (75 ml) and the solution  
10 was cooled in an ice-bath to 0-5°C. Trifluoroacetic acid (37.5 ml) was added followed by the dropwise addition over 15 minutes of a solution of fuming nitric acid (24M; 7.42 ml) in methylene chloride (15 ml). The resultant solution was allowed to warm to ambient temperature and was stirred for 2 hours. Volatile materials were evaporated. The residue was dissolved in methylene chloride (50 ml) and the solution was cooled in an ice-bath to 0-5°C.  
15 Diethyl ether was added and the resultant precipitate was collected and dried under vacuum at 50°C. The solid was dissolved in methylene chloride (500 ml) and hydrogen chloride (3M solution in diethyl ether; 30 ml) was added followed by diethyl ether (500 ml). The resultant solid was collected and dried under vacuum at 50°C. There was thus obtained ethyl  
5-methoxy-4-(N-methylpiperidin-4-ylmethoxy)-2-nitrobenzoate (28.4 g); NMR Spectrum:  
20 (DMSO-d<sub>6</sub>) 1.3 (t, 3H), 1.45-1.65 (m, 2H), 1.75-2.1 (m, 3H), 2.75 (s, 3H), 2.9-3.05 (m, 2H), 3.4-3.5 (d, 2H), 3.95 (s, 3H), 4.05 (d, 2H), 4.3 (q, 2H), 7.32 (s, 1H), 7.66 (s, 1H).

A mixture of a portion (3.89 g) of the material so obtained, 10% platinum-on-activated carbon (50% wet, 0.389 g) and methanol (80 ml) was stirred under 1.8 atmospheres pressure of hydrogen until uptake of hydrogen ceased. The mixture was filtered and the filtrate was  
25 evaporated. The residue was dissolved in water (30 ml) and basified to pH10 by the addition of a saturated aqueous sodium bicarbonate solution. The mixture was diluted with a 1:1 mixture of ethyl acetate and diethyl ether and the organic layer was separated. The aqueous layer was further extracted with a 1:1 mixture of ethyl acetate and diethyl ether and the organic extracts were combined, washed in turn with water and brine, dried over magnesium  
30 sulphate and evaporated. The residue was triturated under a mixture of petroleum ether (b.p. 60-80°C) and diethyl ether. The solid so obtained was isolated, washed with petroleum ether and dried under vacuum at 60°C. There was thus obtained ethyl 2-amino-5-methoxy-4-(N-methylpiperidin-4-ylmethoxy)benzoate (2.58 g); m.p. 111-112°C; NMR Spectrum:

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(CDCl<sub>3</sub>) 1.35 (t, 3H), 1.4-1.5 (m, 2H), 1.85 (m, 3H), 1.95 (t, 2H), 2.29 (s, 3H), 2.9 (d, 2H), 3.8 (s, 3H), 3.85 (d, 2H), 4.3 (q, 2H), 5.55 (br s, 2H), 6.13 (s, 1H), 7.33 (s, 1H).

A mixture of ethyl 2-amino-5-methoxy-4-(N-methylpiperidin-4-ylmethoxy)benzoate (16.1 g), formamidine acetic acid salt (5.2 g) and 2-methoxyethanol (160 ml) was stirred and heated at 115°C for 2 hours. Further formamidine acetic acid salt (10.4 g) was added in portions every 30 minutes during 4 hours and heating was continued for 30 minutes after the last addition. The resultant mixture was evaporated. The solid residue was stirred under a mixture of methylene chloride (50ml) and ethanol (100ml). The precipitate was removed by filtration and the filtrate was concentrated to a final volume of 100ml. The resultant suspension was cooled to 5°C. The solid so obtained was collected, washed with cold ethanol and with diethyl ether and dried under vacuum at 60°C. There was thus obtained 6-methoxy-7-(N-methylpiperidin-4-ylmethoxy)-3,4-dihydroquinazolin-4-one (12.7 g); NMR Spectrum: (DMSO-d<sub>6</sub>) 1.25-1.4 (m, 2H), 1.75 (d, 2H), 1.9 (t, 1H), 1.9 (s, 3H), 2.16 (s, 2H), 2.8 (d, 2H), 3.9 (s, 3H), 4.0 (d, 2H), 7.11 (s, 1H), 7.44 (s, 1H), 7.97 (s, 1H).

A mixture of a portion (2.8 g) of the material so obtained, thionyl chloride (28 ml) and DMF (0.28 ml) was heated to reflux for 1 hour. The mixture was evaporated and the precipitate was triturated under diethyl ether. The resultant solid was isolated and washed with diethyl ether. The solid was then dissolved in methylene chloride and the solution was washed with a saturated aqueous sodium bicarbonate solution. The organic layer was washed in turn with water and brine, dried over magnesium sulphate and evaporated. There was thus obtained 4-chloro-6-methoxy-7-(N-methylpiperidin-4-ylmethoxy)quinazoline (2.9 g), NMR Spectrum: (DMSO-d<sub>6</sub>) 1.3-1.5 (m, 2H), 1.75-1.9 (m, 4H), 2.0 (t, 1H), 2.25 (s, 3H), 2.85 (d, 2H), 4.02 (s, 3H), 4.12 (d, 2H), 7.41 (s, 1H), 7.46 (s, 1H), 8.9 (s, 1H).

**Example 2    4-(2-chloro-5-methoxyanilino)-6-methoxy-7-piperidin-4-ylmethoxyquinazoline monohydrochloride salt**

A mixture of 4-chloro-6-methoxy-7-(N-tert-butoxycarbonylpiperidin-4-ylmethoxy)quinazoline (0.08 g), 2-chloro-5-methoxyaniline hydrochloride (0.042 g), a 6M solution of hydrogen chloride in isopropanol (0.036 ml) and isopropanol (4 ml) was stirred and heated to 80°C for 1.5 hours. The mixture was cooled to ambient temperature and the precipitate was isolated, washed in turn with isopropanol and diethyl ether and dried under vacuum. There was thus obtained the title compound (0.045 g); NMR Spectrum: (DMSO-d<sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.5-1.65 (m, 2H), 1.98 (d, 2H), 2.15-2.3 (m, 1H), 2.95 (t, 2H), 3.35 (d, 2H),

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3.8 (s, 3H), 4.0 (s, 3H), 4.11 (d, 2H), 7.05 (m, 1H), 7.17 (d, 1H), 7.36 (s, 1H), 7.54 (d, 1H), 8.13 (s, 1H), 8.82 (s, 1H); Mass Spectrum: M-H<sup>+</sup> 427 and 429.

The 7-(N-tert-butoxycarbonylpiperidin-4-ylmethoxy)-4-chloro-6-methoxyquinazoline used as a starting material was prepared as follows :-

5 Sodium hydride (60% suspension in mineral oil, 1.44 g) was added portionwise during 20 minutes to a solution of 7-benzyloxy-6-methoxy-3,4-dihydroquinazolin-4-one (International Patent Application WO 97/22596, Example 1 thereof; 8.46 g) in DMF (70 ml). The mixture was stirred at ambient temperature for 1.5 hours. Chloromethyl pivalate (5.65 g) was added dropwise and the mixture was stirred at ambient temperature for 2 hours. The  
10 mixture was diluted with ethyl acetate (100 ml) and poured onto a mixture (400 ml) of ice and water containing 2N aqueous hydrochloric acid (4 ml). The organic layer was separated and the aqueous layer was extracted with ethyl acetate. The combined extracts were washed with brine, dried over magnesium sulphate and evaporated. The residue was triturated under a mixture of diethyl ether and petroleum ether (b.p. 60-80°C) and the resultant solid was  
15 collected and dried under vacuum. There was thus obtained 7-benzyloxy-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one (10 g); NMR Spectrum: (DMSO-d<sub>6</sub>) 1.11 (s, 9H), 3.89 (s, 3H), 5.3 (s, 2H), 5.9 (s, 2H), 7.27 (s, 1H), 7.35 (m, 1H), 7.47 (t, 2H), 7.49 (d, 2H), 7.51 (s, 1H), 8.34 (s, 1H).

A mixture of a portion (7 g) of the material so obtained, 10% palladium-on-charcoal  
20 catalyst (0.7 g), DMF (50 ml), methanol (50 ml), acetic acid (0.7 ml) and ethyl acetate (250 ml) was stirred under an atmosphere pressure of hydrogen for 40 minutes. The catalyst was removed by filtration and the solvent was evaporated. The residue was triturated under diethyl ether and the resultant solid was collected and dried under vacuum. There was thus obtained 7-hydroxy-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one (4.36 g);  
25 NMR Spectrum: (DMSO-d<sub>6</sub>) 1.1 (s, 9H), 3.89 (s, 3H), 5.89 (s, 2H), 7.0 (s, 1H), 7.48 (s, 1H), 8.5 (s, 1H).

Using an analogous procedure to that described in the fourth paragraph of the portion of Example 1 that is concerned with the preparation of starting materials, 7-hydroxy-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one was reacted with  
30 N-tert-butoxycarbonyl-4-(4-toluenesulphonyloxymethyl)piperidine to give 7-(N-tert-butoxycarbonylpiperidin-4-ylmethoxy)-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one.



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A mixture of 7-(N-tert-butoxycarbonylpiperidin-4-ylmethoxy)-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one (6 g) and a saturated methanolic ammonia solution (100ml) was stirred at ambient temperature for 16 hours. The resultant mixture was evaporated and the residue was triturated under diethyl ether. The solid so obtained was isolated, washed with a 49:1 mixture of diethyl ether and methylene chloride and dried under vacuum. There was thus obtained 7-(N-tert-butoxycarbonylpiperidin-4-ylmethoxy)-6-methoxy-3,4-dihydroquinazolin-4-one (3.3 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 1.12-1.3 (m, 2H), 1.42 (s, 9H), 1.8 (d, 2H), 2.02 (m, 1H), 2.7-2.9 (m, 2H), 3.9 (s, 3H), 4.02 (d, 4H), 7.15 (s, 1H), 7.45 (s, 1H), 8.0 (s, 1H).

10 A mixture of a portion (0.2 g) of the material so obtained, carbon tetrachloride (0.15 ml), triphenylphosphine (0.25 g) and 1,2-dichloroethane (10 ml) was stirred and heated to 70°C for 2 hours. The mixture was evaporated and the residue was purified by column chromatography on silica using a 5:4:1 mixture of methylene chloride, ethyl acetate and methanol as eluent. There was thus obtained 7-(N-tert-butoxycarbonylpiperidin-

15 4-ylmethoxy)-4-chloro-6-methoxyquinazoline (0.07 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 1.15-1.3 (m, 2H), 1.45 (s, 9H), 1.8 (d, 2H), 2.08 (m, 1H), 2.7-2.9 (m, 2H), 4.02 (m, 5H), 4.12 (d, 2H), 7.42 (s, 1H), 7.5 (s, 1H), 8.9 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 408.

**Example 3** 4-(2-bromo-5-methoxyanilino)-6-methoxy-7-[2-(N-methylpiperidin-4-yl)ethoxy]quinazoline dihydrochloride salt

A mixture of 4-chloro-6-methoxy-7-[2-(N-methylpiperidin-4-yl)ethoxy]quinazoline (International Patent Application WO 00/47212, example 241; 0.15 g), 2-bromo-5-methoxyaniline (0.113 g), a 6M solution of hydrogen chloride in isopropanol (0.075 ml) and isopropanol (5 ml) was stirred and heated to 80°C for 2 hours. The mixture was cooled to ambient temperature and diethyl ether was added. The resultant solid was isolated, washed with diethyl ether and dried under vacuum. There was thus obtained the title compound (0.12 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.5 (m, 2H), 1.8 (m, 3H), 2.0 (m, 2H), 2.75 (s, 3H), 2.95 (t, 2H), 3.4 (m, 2H), 3.8 (s, 3H), 4.0 (s, 3H), 4.3 (m, 2H), 7.0 (m, 1H), 7.2 (d, 1H), 7.45 (s, 1H), 7.7 (d, 1H), 8.2 (s, 1H), 8.85 (s, 1H); Mass Spectrum: M-H<sup>-</sup> 499 and

30 501.

The 2-bromo-5-methoxyaniline used as a starting material was obtained as follows :-

A mixture of hydrazine hydrate (1 ml), Raney nickel (0.13 g) and methanol was stirred and heated to reflux and a solution of 2-bromo-5-methoxy-1-nitrobenzene (1 g) in methanol

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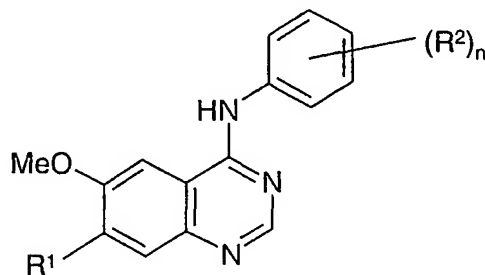
(18 ml) was added dropwise. The resultant mixture was heated to reflux for a further 15 minutes. The reaction mixture was cooled to ambient temperature, filtered and evaporated. The residue was partitioned between methylene chloride and water. The organic phase was dried over magnesium sulphate and evaporated to give 2-bromo-5-methoxyaniline (0.8 g);

5 NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 3.65 (s, 3H), 5.25 (br s, 2H), 6.1 (m, 1H), 6.4 (d, 1H), 7.2 (d, 1H).

**Example 4**

Using an analogous procedure to that described in Example 2 or Example 3, the appropriate 4-chloroquinazoline was reacted with the appropriate aniline to give the

10 compounds described in Table I. Unless otherwise stated, each compound described in Table I was obtained as a dihydrochloride salt.

**Table I**

15

| Compound No. & Note | R <sup>1</sup>                                     | (R <sup>2</sup> ) <sub>n</sub> |
|---------------------|--|--------------------------------|
| [1]                 | 3-pyrrolidin-1-ylpropoxy                           | 2-chloro-5-methoxy             |
| [2]                 | 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy | 2-chloro-5-methoxy             |
| [3]                 | 2-(N-methylpiperidin-4-yl)ethoxy                   | 2-chloro-5-methoxy             |
| [4]                 | 2-piperidin-4-ylethoxy                             | 2-bromo-5-methoxy              |
| [5]                 | piperidin-4-ylmethoxy                              | 2-bromo-5-methoxy              |
| [6]                 | 2-acetoxy-3-piperidinopropoxy                      | 2-bromo-5-methoxy              |
| [7]                 | 2-acetoxy-3-(4-cyanomethylpiperazin-1-yl)propoxy   | 2-bromo-5-methoxy              |

**Notes**

[1] The procedure of Example 3 was followed. The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.9 (m, 2H), 2.05 (m, 2H), 2.3

20 (m, 2H), 3.05 (m, 2H), 3.35 (m, 2H), 3.6 (m, 2H), 3.8 (s, 3H), 4.0 (s, 3H), 4.35 (m, 2H), 7.05

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(m, 1H), 7.15 (d, 1H), 7.45 (s, 1H), 7.55 (d, 1H), 8.25 (s, 1H), 8.8 (s, 1H); Mass Spectrum:  $M+H^+$  443 and 445.

The 4-chloro-7-(3-pyrrolidin-1-ylpropoxy)-6-methoxyquinazoline used as a starting material was prepared as follows :-

5 A mixture of 4-hydroxy-3-methoxybenzoic acid (8.4 g), 3-(pyrrolidin-1-yl)propyl chloride (J. Amer. Chem. Soc., 1955, 77, 2272; 14.75 g), potassium carbonate (13.8 g), potassium iodide (1.66 g) and DMF (150 ml) was stirred and heated to 100°C for 3 hours. The mixture was allowed to cool to ambient temperature, filtered and the filtrate was evaporated. The residue was dissolved in ethanol (75 ml), 2N aqueous sodium hydroxide  
10 solution (75 ml) was added and the mixture was heated to 90°C for 2 hours. The mixture was concentrated by evaporation and acidified by the addition of concentrated aqueous hydrochloric acid. The resultant mixture was washed with diethyl ether and then purified by column chromatography using a Diaion (trade mark of Mitsubishi) HP20SS resin column, eluting with water and then with a gradient of methanol (0 to 25%) in dilute hydrochloric acid  
15 (pH2.2). The methanol was removed by evaporation and the aqueous residue was freeze dried to give 3-methoxy-4-(3-pyrrolidin-1-ylpropoxy)benzoic acid hydrochloride (12.2 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 2.2 (m, 2H), 3.15 (t, 2H), 3.3 (t, 2H), 3.5 (d, 2H), 3.7 (t, 2H), 3.82 (s, 3H), 4.05 (d, 2H), 4.15 (t, 2H), 7.07 (d, 1H), 7.48 (s, 1H), 7.59 (d, 1H).

The material so obtained was dissolved in trifluoroacetic acid (40 ml) and the solution  
20 was cooled to 0°C. Fuming nitric acid (2.4 ml) was added slowly. The cooling bath was removed and the reaction mixture was stirred at ambient temperature for 1 hour. The mixture was evaporated and a mixture of ice and water was added to the residue. The mixture was evaporated. The solid residue was dissolved in dilute hydrochloric acid (pH2.2) and purified by column chromatography using a Diaion HP20SS resin column using a gradient of methanol  
25 (0 to 50%) in water. Concentration of the fractions by evaporation gave a precipitate which was collected and dried under vacuum over phosphorus pentoxide. There was thus obtained 5-methoxy-2-nitro-4-(3-pyrrolidin-1-ylpropoxy)benzoic acid hydrochloride (12.1 g, 90%); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.8-1.9 (m, 2H), 2.0-2.1 (m, 2H), 2.1-2.2 (m, 2H), 3.0-3.1 (m, 2H), 3.3 (t, 2H), 3.6-3.7 (m, 2H), 3.95 (s, 3H), 4.25 (t, 2H), 7.35 (s, 1H), 7.62 (s,  
30 1H).

A mixture of a portion (9.63 g) of the material so obtained, thionyl chloride (20 ml) and DMF (0.05 ml) was heated to 45°C for 1.5 hours. The excess thionyl chloride was evaporated using the evaporation of added toluene (x2) to remove the last traces. The

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resultant solid was suspended in a mixture of THF (250 ml) and methylene chloride (100ml) and ammonia was bubbled through the mixture for 30 minutes. The resultant mixture was stirred for a further 1.5 hours at ambient temperature. The volatiles were removed by evaporation and the residue was dissolved in water and purified by column chromatography using a Diaion HP20SS resin column eluting with a gradient of methanol (0 to 5%) in water. The solvent was removed by evaporation from the fractions containing product. The residue was dissolved in a minimum of methanol and the solution was diluted with diethyl ether. The resultant precipitate was collected by filtration, washed with diethyl ether and dried under vacuum to give 5-methoxy-2-nitro-4-(3-pyrrolidin-1-ylpropoxy)benzamide (7.23 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.85-1.95 (m, 2H), 2-2.1 (m, 2H), 2.15-2.25 (m, 2H), 3.0-3.1 (m, 2H), 3.31 (t, 2H), 3.62 (t, 2H), 3.93 (s, 3H), 4.2 (t, 2H), 7.16 (s, 1H), 7.6 (s, 1H).

A mixture of a portion (1.5 g) of the material so obtained, concentrated aqueous hydrochloric acid (5 ml) and methanol (20 ml) was warmed to 50°C to give a solution. Iron powder (1.3 g) was added in portions and the reaction mixture was heated to reflux for 1 hour. The mixture was allowed to cool to ambient temperature. Insoluble material was removed by filtration through diatomaceous earth and the filtrate was evaporated. The residue was purified by column chromatography using a Diaion HP20SS resin column, eluting with water and then with dilute aqueous hydrochloric acid (pH2). The fractions containing product were concentrated by evaporation and the resultant precipitate was collected by filtration and dried under vacuum over phosphorus pentoxide. There was thus obtained 2-amino-5-methoxy-4-(3-pyrrolidin-1-ylpropoxy)benzamide hydrochloride (1.44 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.9 (br s, 2H), 2.05 (br s, 2H), 2.2 (br s, 2H), 3.05 (br s, 2H), 3.3 (t, 2H), 3.61 (br s, 2H), 3.8 (s, 3H), 4.11 (t, 2H), 7.05 (s, 1H), 7.53 (s, 1H).

After repetition of the previous reaction, a mixture of 2-amino-5-methoxy-4-(3-pyrrolidin-1-ylpropoxy)benzamide hydrochloride (5.92 g), Gold's reagent (3.5 g) and dioxane (50 ml) was heated to reflux for 5 hours. Acetic acid (0.7 ml) and sodium acetate (1.33 g) were added and the reaction mixture was heated to reflux for a further 5 hours. The mixture was allowed to cool to ambient temperature and evaporated. The residue was dissolved in water, adjusted to pH8 with 2N aqueous sodium hydroxide solution and purified on a Diaion HP20SS resin column eluting with methanol (gradient 0-50 %) in water. The fractions containing product were concentrated by evaporation and then freeze dried to give 6-methoxy-7-(3-pyrrolidin-1-ylpropoxy)-3,4-dihydroquinazolin-4-one (4.55 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.9 (m, 2H), 2.0-2.1 (m, 2H), 2.2-2.3 (m, 2H), 3.05 (m,

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2H), 3.34 (t, 2H), 3.6-3.7 (br s, 2H), 3.94 (s, 3H), 4.27 (t, 2H), 7.31 (s, 1H), 7.55 (s, 1H), 9.02 (s, 1H).

A mixture of a portion (1.7 g) of the material so obtained, thionyl chloride (25 ml) and DMF (0.2 ml) was heated at reflux for 3 hours. Excess thionyl chloride was removed by  
5 evaporation and by azeotrope with toluene (x2). The residue was suspended in diethyl ether and washed with a 10% aqueous solution of sodium bicarbonate. The organic layer was dried over magnesium sulphate and evaporated to give 4-chloro-6-methoxy-7-(3-pyrrolidin-1-ylpropoxy)quinazoline (1.94 g); NMR Spectrum: (CDCl<sub>3</sub>) 1.8 (br s, 4H), 2.17 (m, 2H), 2.6 (br s, 4H), 2.7 (t, 2H), 4.05 (s, 3H), 4.3 (t, 2H), 7.35 (s, 1H), 7.38 (s, 1H), 8.86 (s, 1H).

10 [2] The procedure of Example 3 was followed. The product gave the following characterising data: NMR Spectrum: (DMSO-d<sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 2.35 (m, 2H), 3.5 (m, 2H), 3.7 (m, 4H), 3.8 (s, 3H), 3.85 (m, 4H), 4.0 (s, 3H), 4.35 (m, 2H), 7.05 (m, 1H), 7.2 (d, 1H), 7.4 (s, 1H), 7.6 (d, 1H), 8.2 (s, 1H), 8.85 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 507 and 509.

The 4-chloro-7-[3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy]-  
15 6-methoxyquinazoline used as a starting material was prepared as follows :-

A mixture of 2-amino-4-benzyloxy-5-methoxybenzamide (J. Med. Chem., 1977, 20, 146-149; 10 g), (3-dimethylamino-2-azaprop-2-en-1-ylidene)dimethylammonium chloride (Gold's reagent, 7.4 g) and dioxane (100 ml) was stirred and heated to reflux for 24 hours. Sodium acetate (3.02 g) and acetic acid (1.65 ml) were added and the reaction mixture was  
20 heated for a further 3 hours. The mixture was evaporated and water was added to the residue. The resultant solid was collected by filtration, washed with water and dried. The material was recrystallised from acetic acid to give 7-benzyloxy-6-methoxy-3,4-dihydroquinazolin-4-one (8.7 g).

After repetition of the reaction so described, a mixture of 7-benzyloxy-6-methoxy-  
25 3,4-dihydroquinazolin-4-one (35 g), thionyl chloride (440 ml) and DMF (1.75 ml) was heated to reflux for 4 hours. The thionyl chloride was evaporated under vacuum and the residue was azeotroped with toluene three times. The residue was dissolved in N-methylpyrrolidin-2-one (250 ml) to give a solution of 7-benzyloxy-4-chloro-6-methoxyquinazoline.

A mixture of 7-benzyloxy-6-methoxy-3,4-dihydroquinazolin-4-one (20.3 g), thionyl  
30 chloride (440 ml) and DMF (1.75 ml) was heated to reflux for 4 hours. The thionyl chloride was evaporated under vacuum and the residue was azeotroped with toluene three times to give 7-benzyloxy-4-chloro-6-methoxyquinazoline.

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A mixture of the 7-benzyloxy-4-chloro-6-methoxyquinazoline so obtained, potassium carbonate (50 g) and 4-chloro-2-fluorophenol (8.8 ml) and DMF (500 ml) was stirred and heated to 100°C for 5 hours. The mixture was allowed to cool to ambient temperature, poured into water (2 L) and stirred at ambient temperature for a few minutes. The resultant solid was isolated and washed with water. The solid was dissolved in methylene chloride and the solution was filtered and treated with decolourising charcoal. The resultant solution was filtered and evaporated to give a solid which was triturated under diethyl ether. There was thus obtained 7-benzyloxy-4-(4-chloro-2-fluorophenoxy)-6-methoxyquinazoline (23.2 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 3.98 (s, 3H), 5.34 (s, 2H), 7.42 (m, 9H), 7.69 (m, 1H), 8.55 (s, 1H).

A mixture of the material so obtained and trifluoroacetic acid (15 ml) was heated to reflux for 3 hours. The reaction mixture was allowed to cool, toluene was added and the mixture was evaporated. The residue was triturated under diethyl ether and then under acetone. The resultant precipitate was isolated and dried to give 4-(4-chloro-2-fluorophenoxy)-7-hydroxy-6-methoxyquinazoline trifluoroacetate salt (21.8 g) which was used without further purification.

3-(1,1-Dioxotetrahydro-4H-1,4-thiazin-4-yl)propan-1-ol (4.2 g) and 1,1'-(azodicarbonyl)dipiperidine (11.7 g) were added in turn to a mixture of 4-(4-chloro-2-fluorophenoxy)-7-hydroxy-6-methoxyquinazoline (5.0 g), tributylphosphine (11.1 ml) and methylene chloride (150 ml). The resultant mixture was stirred at ambient temperature overnight. The mixture was diluted with diethyl ether (300 ml) and the precipitate was removed by filtration. The filtrate was evaporated and the residue was purified by column chromatography on silica using a 19:1 mixture of methylene chloride and methanol as eluent. The material so obtained was triturated under ethyl acetate and dried to give 4-(4-chloro-2-fluorophenoxy)-7-[3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy]-6-methoxyquinazoline (5.4 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 1.86 (m, 2H), 2.65 (t, 2H), 2.92 (m, 4H), 3.08 (m, 4H), 3.97 (s, 3H), 4.26 (t, 2H), 7.4 (m, 1H), 7.42 (s, 1H), 7.56 (m, 2H), 7.68 (m, 1H), 8.54 (s, 1H).

A mixture of a portion (3.5 g) of the material so obtained and a 2N aqueous hydrochloric acid solution (56 ml) was stirred and heated to 95°C for 2 hours. The reaction mixture was cooled to ambient temperature and treated with solid sodium bicarbonate to give a thick paste which was diluted with water and filtered. The solid was transferred to a flask and azeotroped with toluene twice to give a dry solid. The solid was purified by column

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chromatography on silica using a 19:1 mixture of methylene chloride and methanol as eluent. There was thus obtained 7-[3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy]-6-methoxy-3,4-dihydroquinazolin-4-one (2.26 g) as a white solid; Mass Spectrum:  $M+H^+$  368.

After repetition of the previous reaction, a mixture of 7-[3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy]-6-methoxy-3,4-dihydroquinazolin-4-one (4.2 g), thionyl chloride (45 ml) and DMF (0.1 ml) was heated to reflux for 2.5 hours. The residue was diluted with toluene and was evaporated under vacuum. The residue was taken up in water and basified to pH8 with a saturated aqueous sodium bicarbonate solution. The mixture was extracted with methylene chloride and the organic layer was washed in turn with water and brine. The organic solution was filtered through phase separating paper and evaporated to give an orange solid which was purified by column chromatography on silica using a 19:1 mixture of methylene chloride and methanol as eluent. The resultant solid was triturated under diethyl ether and dried to give 4-chloro-7-[3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy]-6-methoxyquinazoline (2.27 g); Mass Spectrum:  $M+H^+$  386.

The 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propan-1-ol used as an intermediate was obtained as follows :-

A mixture of 3-aminopropan-1-ol (0.65 ml) and divinyl sulphone (1 g) was heated to 110°C for 45 minutes. The mixture was allowed to cool to ambient temperature and was purified by column chromatography on silica using a 19:1 mixture of methylene chloride and methanol as eluent. There was thus obtained 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propan-1-ol (0.8 g); NMR Spectrum: ( $CDCl_3$ ) 1.7-1.8 (m, 2H), 2.73 (t, 2H), 3.06 (br s, 8H), 3.25 (s, 1H), 3.78 (t, 2H); Mass Spectrum:  $M+H^+$  194.

[3] The procedure of Example 3 was followed. The reaction product so obtained was mixed with methylene chloride (5 ml) and a saturated methanolic ammonia solution (0.5 ml) was added. The mixture was stirred at ambient temperature for 10 minutes. The mixture was filtered and the filtrate was evaporated. The residue was purified by column chromatography on silica using a 3:1 mixture of methylene chloride and a saturated methanolic ammonia solution as eluent. The material so obtained was dissolved in diethyl ether and a 2.9M solution of hydrogen chloride in diethyl ether (0.5 ml) was added. The mixture was evaporated and the residue was triturated under diethyl ether. The resultant solid was isolated, washed with diethyl ether and dried under vacuum. There was thus obtained the dihydrochloride salt of the required compound which gave the following characterising data: NMR Spectrum: ( $DMSO-d_6$ ) 1.4-1.5 (m, 2H), 1.7-1.8 (m, 3H), 2.0 (d, 2H), 2.75 (s, 3H), 2.95

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(m, 2H), 3.4 (m, 2H), 3.8 (s, 3H), 3.95 (s, 3H), 4.2 (m, 2H), 6.92 (m, 1H), 7.12 (d, 1H), 7.22 (s, 1H), 7.5 (d, 1H), 7.85 (s, 1H), 8.32 (s, 1H), 9.6 (m, 1H); Mass Spectrum:  $M+H^+$  457 and 459.

[4] 7-[2-(N-tert-Butoxycarbonylpiperidin-4-yl)ethoxy]-4-chloro-6-methoxyquinazoline  
5 was used as the appropriate 4-chloroquinazoline and the procedure of Example 2 was followed. The product was obtained as the monohydrochloride salt and gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.88 (br s, 3H), 2.0 (d, 2H), 2.95 (m, 2H), 3.32 (d, 2H), 3.83 (s, 3H), 4.02 (s, 3H), 4.3 (m, 2H), 7.02 (m, 1H), 7.20 (d, 1H), 7.38 (s, 1H), 7.72 (d, 1H), 8.16 (s, 1H), 8.85 (s, 1H); Mass Spectrum:  $M-H^-$  485 and 487.

10 The 7-[2-(N-tert-butoxycarbonylpiperidin-4-yl)ethoxy]-4-chloro-6-methoxyquinazoline used as a starting material was obtained as follows :-

A mixture of 7-hydroxy-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one (2 g), N-tert-butoxycarbonyl-4-[2-(4-toluenesulphonyloxy)ethyl]piperidine (2.84 g), potassium carbonate (1.8 g) and DMF (20 ml) was stirred and heated to 95°C for 2.5 hours. The  
15 resultant mixture was cooled to ambient temperature and poured onto a mixture of ice and water. The mixture was extracted with methylene chloride. The organic layer was washed with brine, dried over magnesium sulphate and evaporated. There was thus obtained 7-[2-(N-tert-butoxycarbonylpiperidin-4-yl)ethoxy]-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one (2 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 1.0-1.15 (m, 2H), 1.15 (s, 9H), 1.4 (s, 9H), 1.6-1.8 (m, 3H), 2.6-2.8 (m, 2H), 3.92 (s, 3H), 3.9-4.0 (m, 2H), 4.2 (m, 2H),  
20 5.92 (s, 2H), 7.2 (s, 1H), 7.5 (s, 1H), 8.3 (s, 1H).

Using an analogous procedure to that described in the fourth paragraph of the portion of Example 2 that is concerned with the preparation of starting materials,

7-[2-(N-tert-butoxycarbonylpiperidin-4-yl)ethoxy]-6-methoxy-3-pivaloyloxymethyl-  
25 3,4-dihydroquinazolin-4-one (2 g) was treated with a saturated methanolic ammonia solution to give 7-[2-(N-tert-butoxycarbonylpiperidin-4-yl)ethoxy]-6-methoxy-3,4-dihydroquinazolin-4-one (1.3 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 1.0-1.15 (m, 2H), 1.4 (s, 9H), 1.6-1.8 (m, 3H), 2.6-2.8 (m, 2H), 3.3-3.5 (m, 2H), 3.9 (s, 3H), 3.9-4.0 (m, 2H), 4.18 (m, 2H), 7.15 (s, 1H), 7.45 (s, 1H), 8.0 (s, 1H); Mass Spectrum:  $M+H^+$  404.

30 Using an analogous procedure to that described in the fifth paragraph of the portion of Example 2 that is concerned with the preparation of starting materials,

7-[2-(N-tert-butoxycarbonylpiperidin-4-yl)ethoxy]-6-methoxy-3,4-dihydroquinazolin-4-one (0.2 g) was reacted with carbon tetrachloride and triphenylphosphine to give



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7-[2-(N-tert-butoxycarbonylpiperidin-4-yl)ethoxy]-4-chloro-6-methoxyquinazoline (0.03 g);  
NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 1.0-1.2 (m, 2H), 1.4 (s, 9H), 1.6-1.8 (m, 5H), 2.6-2.8 (m, 2H),  
3.92 (d, 2H), 4.0 (s, 3H), 4.3 (m, 2H), 7.4 (s, 1H), 7.5 (s, 1H), 8.9 (s, 1H).

The N-tert-butoxycarbonyl-4-[2-(4-toluenesulphonyloxy)ethyl]piperidine used as a  
5 starting material was prepared by the reaction of 4-toluenesulphonyl chloride with  
N-tert-butoxycarbonyl-4-(2-hydroxyethyl)piperidine (International Patent Application  
WO 00/47212, in example 126 thereof) using an analogous procedure to that described in the  
third paragraph of the portion of Example 1 that is concerned with the preparation of starting  
materials.

10 [5] 7-(N-tert-Butoxycarbonylpiperidin-4-ylmethoxy)-4-chloro-6-methoxyquinazoline was  
used as the appropriate 4-chloroquinazoline and the procedure of Example 2 was followed.  
The product was obtained as the monohydrochloride salt and gave the following characterising  
data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 1.5-1.7 (m, 2H), 1.95-2.05 (m, 2H), 2.15-2.25 (m, 1H), 2.9-  
3.05 (m, 2H), 3.3-3.4 (m, 2H), 3.8 (s, 3H), 4.05 (s, 3H), 4.12 (d, 2H), 7.03 (m, 1H), 7.18 (d,  
15 1H), 7.45 (s, 1H), 7.72 (d, 1H), 8.28 (s, 1H), 8.7 (br s, 1H), 8.8 (s, 1H), 9.05 (br s, 1H); Mass  
Spectrum: M+H<sup>+</sup> 473 and 475.

[6] The procedure of Example 3 was followed. The product gave the following  
characterising data: Mass Spectrum: M+H<sup>+</sup> 559 and 561.

The 7-(2-acetoxy-3-piperidinopropoxy)-4-chloro-6-methoxyquinazoline used as a  
20 starting material was prepared as follows :-

A mixture of 7-hydroxy-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one  
(40 g), 2,3-epoxypropyl bromide (16.8 ml), potassium carbonate (36 g) and DMF (400 ml)  
was stirred and heated to 70°C for 1.5 hours. The mixture was poured into an ice-water  
mixture (1.5 L) and the resultant precipitate was isolated, washed in turn with water and  
25 diethyl ether and dried under vacuum over phosphorus pentoxide. There was thus obtained  
7-(2,3-epoxypropoxy)-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one (46.7 g).

A mixture of a portion (8 g) of the material so obtained, piperidine (2.4 ml) and  
chloroform (120 ml) was heated to reflux for 16 hours. The mixture was evaporated and the  
residue was purified by column chromatography on silica using a 19:1 mixture of methylene  
30 chloride and methanol as eluent. There was thus obtained 7-(2-hydroxy-  
3-piperidinopropoxy)-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one as a solid  
(9.2 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 1.1 (s, 9H), 1.5 (m, 6H), 2.4 (m, 6H), 3.9 (s, 3H), 4.05  
(m, 2H), 4.15 (m, 1H), 4.9 (br s, 1H), 5.9 (s, 2H), 7.2 (s, 1H), 7.5 (s, 1H), 8.35 (s, 1H).

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A mixture of the material so obtained and a saturated methanolic ammonia solution (240 ml) was stirred at ambient temperature for 48 hours. The mixture was evaporated and the resultant solid was washed with a 19:1 mixture of diethyl ether and methylene chloride. There was thus obtained 7-(2-hydroxy-3-piperidinopropoxy)-6-methoxy-

5 3,4-dihydroquinazolin-4-one (6.2 g); NMR Spectrum: (DMSO-d<sub>6</sub> & CF<sub>3</sub>CO<sub>2</sub>D) 1.4 (m, 1H), 1.7 (m, 2H), 1.8 (m, 3H), 3.0 (m, 2H), 3.35 (m, 2H), 3.5 (m, 2H), 3.9 (s, 3H), 4.15 (d, 2H), 4.4 (m, 1H), 7.3 (s, 1H), 7.55 (s, 1H), 8.75 (s, 1H).

A mixture of a portion (5.6 g) of the material so obtained and acetic anhydride (8.25 ml) was stirred at ambient temperature for 1.3 hours. Water (3 ml) was added and the  
10 resultant mixture was cooled in an ice-bath and basified to pH9.5 by the addition of a 2N aqueous sodium hydroxide solution. The mixture was extracted with methylene chloride and the organic phase was washed with water and with brine, dried over magnesium sulphate and evaporated. There was thus obtained 7-(2-acetoxy-3-piperidinopropoxy)-6-methoxy-3,4-dihydroquinazolin-4-one as a solid (4.8 g) which was used without further purification.

15 A mixture of the material so obtained, thionyl chloride (62 ml) and DMF (0.7 ml) was heated to reflux for 1.5 hours. The mixture was evaporated, toluene was added and the mixture was evaporated. Methylene chloride followed by a mixture of ice and water were added to the residue and the mixture was basified to pH7.5 by the addition of a saturated aqueous sodium bicarbonate solution and to pH9 by the addition of a 2N aqueous sodium  
20 hydroxide solution. The organic phase was washed with water and with brine, dried over magnesium sulphate and evaporated. The residue was triturated under diethyl ether. There was thus obtained 7-(2-acetoxy-3-piperidinopropoxy)-4-chloro-6-methoxyquinazoline; NMR Spectrum: (CDCl<sub>3</sub> and CD<sub>3</sub>CO<sub>2</sub>D) 1.6 (m, 2H), 1.9 (m, 4H), 2.1 (s, 3H), 3.2 (br s, 4H), 3.5 (m, 2H), 4.05 (s, 3H), 4.35 (m, 2H), 5.7 (m, 1H), 7.4 (s, 1H), 7.5 (s, 1H), 8.9 (s, 1H).

25 [7] The procedure of Example 3 was followed. The product gave the following characterising data: Mass Spectrum: M+H<sup>+</sup> 599 and 601.

The 7-[2-acetoxy-3-(4-cyanomethylpiperazin-1-yl)propoxy]-4-chloro-6-methoxyquinazoline used as a starting material was prepared as follows :-

7-(2,3-Epoxypropoxy)-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one  
30 was reacted with 1-cyanomethylpiperazine using an analogous procedure to that described in the second paragraph of the portion of Note [6] immediately above that is concerned with the preparation of starting materials. There was thus obtained 7-[3-(4-cyanomethylpiperazin-1-yl)-2-hydroxypropoxy]-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one.

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The material so obtained was taken through an analogous sequence of reactions to those described in the third to fifth paragraphs of the portion of Note [6] immediately above that is concerned with the preparation of starting materials. There was thus obtained

7-[2-acetoxy-3-(4-cyanomethylpiperazin-1-yl)propoxy]-4-chloro-6-methoxyquinazoline;

5 NMR Spectrum: (CDCl<sub>3</sub>) 2.1 (s, 3H), 2.65 (br s, 10H), 3.5 (s, 2H), 4.05 (s, 3H), 4.4 (m, 2H), 5.45 (m, 1H), 7.25 (s, 1H), 7.4 (s, 1H), 8.85 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 434 and 436.

The 1-cyanomethylpiperazine used as a starting material was prepared as follows :-

A mixture of 1-(tert-butoxycarbonyl)piperazine (5 g), 2-chloroacetonitrile (1.9 ml), potassium carbonate (4 g) and DMF (20 ml) was stirred at ambient temperature for 16 hours.

10 A saturated aqueous ammonium chloride solution was added and the mixture was extracted with ethyl acetate. The organic phase was dried over magnesium sulphate and evaporated. The residue was purified by column chromatography on silica using diethyl ether as eluent. There was thus obtained 1-(tert-butoxycarbonyl)-4-cyanomethylpiperazine as a solid (5.7 g); NMR Spectrum: (CDCl<sub>3</sub>) 1.45 (s, 9H), 2.5 (m, 4H), 3.45 (m, 4H), 3.55 (s, 2H).

15 A mixture of the material so obtained, trifluoroacetic acid (20 ml) and methylene chloride (25 ml) was stirred at ambient temperature for 4 hours. The mixture was evaporated, toluene was added and the mixture was evaporated again. The residue was purified by column chromatography on silica using a 9:1 mixture of methylene chloride and methanol as eluent. There was thus obtained 1-cyanomethylpiperazine trifluoroacetate salt which was  
20 treated with solid sodium bicarbonate in a mixture of methylene chloride, ethyl acetate and methanol to give the free base form (2.9 g); NMR Spectrum: (CDCl<sub>3</sub> and DMSO-d<sub>6</sub>) 2.7 (m, 4H), 3.2 (m, 4H), 3.6 (s, 2H), 6.2 (br s, 1H).

#### Example 5 4-(2-chloro-5-methoxyanilino)-6-methoxy-

25 7-[2-(4-pyridyloxy)ethoxy]quinazoline

Diethyl azodicarboxylate (0.142 ml) was added dropwise to a stirred mixture of 4-(2-chloro-5-methoxyanilino)-7-hydroxy-6-methoxyquinazoline (0.2 g), 2-(4-pyridyloxy)ethanol (0.08 g), triphenylphosphine (0.19 g) and methylene chloride (8 ml) and the reaction mixture was stirred at ambient temperature for 1 hour. The mixture was  
30 evaporated and the residue was purified by column chromatography on silica using a 19:1 mixture of methylene chloride and methanol as eluent. There was thus obtained the title compound (0.158 g); NMR Spectrum: (DMSO-d<sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 3.8 (s, 3H), 4.0 (s, 3H), 4.7

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(m, 2H), 4.9 (m, 2H), 7.1 (m, 1H), 7.2 (d, 1H), 7.45 (s, 1H), 7.6 (d, 1H), 7.7 (d, 2H), 8.15 (s, 1H), 8.85 (m, 3H); Mass Spectrum:  $M+H^+$  453 and 455.

The 4-(2-chloro-5-methoxyanilino)-7-hydroxy-6-methoxyquinazoline used as a starting material was obtained as follows :-

5 A mixture of 7-benzyloxy-6-methoxy-3,4-dihydroquinazolin-4-one (35 g), thionyl chloride (440 ml) and DMF (1.75 ml) was heated to reflux for 4 hours. The thionyl chloride was evaporated under vacuum and the residue was azeotroped with toluene three times. The residue was dissolved in N-methylpyrrolidin-2-one (250 ml) to give a solution of 7-benzyloxy-4-chloro-6-methoxyquinazoline.

10 A mixture of 7-benzyloxy-4-chloro-6-methoxyquinazoline (4.3 g), 2-chloro-5-methoxyaniline (2.7 g), a 6.2M solution of hydrogen chloride in isopropanol (0.225 ml) and isopropanol (200 ml) was stirred and heated to 80°C for 2.5 hours. The mixture was cooled to 0°C and the precipitate was isolated, washed with in turn with isopropanol and diethyl ether and dried under vacuum. There was thus obtained 7-benzyloxy-4-(2-chloro-  
15 5-methoxyanilino)-6-methoxyquinazoline (4.73 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 3.8 (s, 3H), 4.03 (s, 3H), 5.36 (s, 2H), 7.06 (m, 1H), 7.18 (d, 1H), 7.4-7.6 (m, 7H), 8.2 (s, 1H), 8.77 (s, 1H), 11.5 (br s, 1H); Mass Spectrum:  $M+H^+$  422 and 424.

A mixture of the material so obtained and trifluoroacetic acid (40 ml) was stirred and heated to 80°C for 4 hours. The mixture was poured into water and solid sodium bicarbonate  
20 was added to basify the mixture to pH8. The resultant precipitate was isolated, washed with water and dried under vacuum at 50°C for 48 hours. The material so obtained was purified by column chromatography on silica using a 1:1 mixture of methylene chloride and ethyl acetate as eluent. There was thus obtained 4-(2-chloro-5-methoxyanilino)-7-hydroxy-  
6-methoxyquinazoline (2.9 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 3.8 (s, 3H), 4.0 (s, 3H), 6.95 (m,  
25 1H), 7.1 (s, 1H), 7.15 (s, 1H), 7.5 (d, 1H), 7.8 (s, 1H), 8.3 (s, 1H), 9.5 (br s, 1H), 10.4 (br s, 1H).

The 2-(4-pyridyloxy)ethanol used as a starting material was prepared using an analogous procedure to that described in J. Chem. Soc. Perkin II, 1987, 1867.

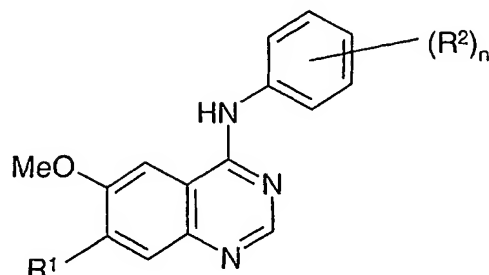
### 30 Example 6

Using an analogous procedure to that described in Example 5, the appropriate 7-hydroxyquinazoline was reacted with the appropriate alcohol to give the compounds

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described in Table II. Unless otherwise stated, each compound described in Table II was obtained as a free base.

Table II



5

| Compound No. & Note | R <sup>1</sup>                                     | (R <sup>2</sup> ) <sub>n</sub> |
|---------------------|--|--------------------------------|
| [1]                 | 3-(4-pyridyloxy)propoxy                            | 2-chloro-5-methoxy             |
| [2]                 | 2-(4-pyridyloxy)ethoxy                             | 2-bromo-5-methoxy              |
| [3]                 | 3-(4-pyridyloxy)propoxy                            | 2-bromo-5-methoxy              |
| [4]                 | 3-(2-pyridyloxy)propoxy                            | 2-chloro-5-methoxy             |
| [5]                 | 2-cyanopyrid-4-ylmethoxy                           | 2-chloro-5-methoxy             |
| [6]                 | 2-cyanopyrid-4-ylmethoxy                           | 2-bromo-5-methoxy              |
| [7]                 | 2-(5-methyl-2-morpholinomethylimidazol-1-yl)ethoxy | 2-chloro-5-methoxy             |

### Notes

[1] The product gave the following characterising data: NMR Spectrum: (DMSO-d<sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 2.4 (m, 2H), 3.8 (s, 3H), 4.0 (s, 3H), 4.4 (m, 2H), 4.6 (m, 2H), 7.1 (m, 1H), 7.2 (d, 1H), 7.4 (s, 1H), 7.6 (m, 3H), 8.1 (s, 1H), 8.8 (d, 2H), 8.85 (s, 1H); Mass Spectrum: M-H<sup>+</sup> 465 and 467.

The 3-(4-pyridyloxy)propanol used as a starting material was prepared as follows :-

Sodium hydroxide (6.66 g) was added to a stirred mixture of 4-chloropyridine hydrochloride (10 g), 1,3-propanediol (24 ml) and DMSO (100 ml) and the resultant mixture was heated to 100°C for 20 hours. The mixture was evaporated and the residue was poured into an ice-water mixture and extracted with ethyl acetate. The organic solution was dried over magnesium sulphate and evaporated. The material so obtained was purified by column chromatography on silica using increasingly polar solvent mixtures of methylene chloride, ethyl acetate and methanol as eluent. There was thus obtained 3-(4-pyridyloxy)propanol

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(3.13 g); NMR Spectrum: (CDCl<sub>3</sub>) 2.05 (m, 2H), 2.95 (br s, 1H), 3.85 (m, 2H), 4.15 (t, 2H), 6.8 (d, 2H), 8.35 (d, 2H).

[2] The product gave the following characterising data; NMR Spectrum: (DMSOd<sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 3.8 (s, 3H), 4.0 (s, 3H), 4.7 (m, 2H), 4.9 (m, 2H), 7.1 (m, 1H), 7.2 (d, 1H), 7.45 (s, 1H), 7.7 (m, 3H), 8.15 (s, 1H), 8.85 (m, 3H); Mass Spectrum: M+H<sup>+</sup> 497 and 499.

[3] The product gave the following characterising data: NMR Spectrum: (DMSOd<sub>6</sub>) 2.3 (m, 2H), 3.8 (s, 3H), 4.0 (s, 3H), 4.3 (m, 4H), 6.9 (m, 1H), 7.05 (d, 2H), 7.15 (s, 1H), 7.25 (s, 1H), 7.65 (d, 1H), 7.85 (s, 1H), 8.35 (s, 1H), 8.4 (d, 2H), 9.5 (s, 1H); Mass Spectrum: M-H<sup>-</sup> 509 and 511.

10 [4] The product gave the following characterising data: NMR Spectrum: (DMSOd<sub>6</sub>) 2.2 (m, 2H), 3.8 (s, 3H), 3.95 (s, 3H), 4.05 (t, 2H), 4.15 (t, 2H), 6.2 (t, 1H), 6.4 (d, 1H), 6.9 (m, 1H), 7.15 (d, 1H), 7.17 (s, 1H), 7.4 (m, 1H), 7.5 (d, 1H), 7.65 (m, 1H), 7.85 (s, 1H), 8.3 (s, 1H), 9.5 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 467 and 469.

The 3-(2-pyridyloxy)propanol used as a starting material is described in

15 Bull. Soc. Chim. Fr., 1970, 637.

[5] The product gave the following characterising data: NMR Spectrum: (DMSOd<sub>6</sub>) 3.8 (s, 3H), 4.05 (s, 3H), 5.5 (s, 2H), 6.95 (m, 1H), 7.15 (d, 1H), 7.3 (s, 1H), 7.5 (d, 1H), 7.85 (m, 1H), 7.95 (s, 1H), 8.15 (s, 1H), 8.35 (s, 1H), 8.85 (d, 1H), 9.6 (br s, 1H); Mass Spectrum: M+H<sup>+</sup> 448 and 450.

20 The 2-cyano-4-hydroxymethylpyridine used as a starting material was prepared as follows :-

Using an analogous procedure to that disclosed in J. Het. Chem., 1993, 30, 631, 4-hydroxymethylpyridine was converted into 4-(tert-butyldimethylsilyloxymethyl)pyridine-2-carbonitrile.

25 A mixture of the material so obtained (3.37 g) tert-butylammonium fluoride (1M solution in THF; 24 ml) and THF (20 ml) was stirred at ambient temperature for 1 hour. The mixture was evaporated and the residue was partitioned between ethyl acetate and a saturated aqueous ammonium chloride solution. The organic phase was washed with water and with brine, dried over magnesium sulphate and evaporated. The residue was purified by column  
30 chromatography on silica using increasingly polar mixtures of petroleum ether (b.p. 40-60°C) and ethyl acetate as eluent. There was thus obtained 2-cyano-4-hydroxymethylpyridine as a solid (1.37 g); NMR Spectrum: (CDCl<sub>3</sub>) 2.25 (br s, 1H), 4.85 (s, 2H), 7.55 (d, 1H), 7.75 (s, 1H), 8.7 (d, 1H).

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[6] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 3.8 (s, 3H), 4.05 (s, 3H), 5.5 (s, 2H), 6.9 (m, 1H), 7.15 (d, 1H), 7.3 (s, 1H), 7.65 (d, 1H), 7.85 (m, 1H), 7.95 (s, 1H), 8.15 (s, 1H), 8.35 (s, 1H), 8.85 (d, 1H), 9.6 (br s, 1H); Mass Spectrum: M+H<sup>+</sup> 492 and 494.

5 [7] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 2.3 (s, 3H), 2.35 (br s, 4H), 3.55 (br s, 4H), 3.65 (s, 2H), 3.8 (s, 3H), 3.95 (s, 3H), 4.5 (br s, 4H), 6.5 (s, 1H), 6.9 (m, 1H), 7.15 (d, 1H), 7.2 (s, 1H), 7.5 (d, 1H), 7.8 (s, 1H), 8.3 (s, 1H), 9.5 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 539 and 541.

The 1-(2-hydroxyethyl)-5-methyl-2-morpholinomethylimidazole used as a starting  
10 material was prepared as follows :-

A mixture of 4-methyl-1-tritylimidazole (J. Heterocyclic Chem., 1982, 19, 253; 32.5 g), methyl bromoacetate (11.4 ml) and acetone (500 ml) was heated to reflux for 2 hours. The solvent was removed by evaporation and the residue was dissolved in methanol (100 ml) and heated to reflux for 45 minutes. The mixture was evaporated and the residue was  
15 triturated under diethyl ether. The resultant precipitate was isolated and stirred at ambient temperature for 1 hour in a mixture of diethyl ether (200 ml) and a saturated methanolic ammonia solution (20 ml). The mixture was filtered and the filtrate was evaporated. The residue was purified by column chromatography on silica using a 49:1 mixture of methylene chloride and methanol as eluent. There was thus obtained methyl 2-(5-methylimidazol-  
20 1-yl)acetate (6 g); NMR Spectrum: (CDCl<sub>3</sub>) 2.16 (s, 3H), 3.78 (s, 3H), 4.61 (s, 3H), 6.8 (s, 1H), 7.42 (s, 1H).

A solution of a portion (1.7 g) of the material so obtained in diethyl ether (20 ml) was added dropwise to a stirred suspension of lithium aluminium hydride (0.76 g) in diethyl ether (70 ml) which was cooled to 0°C. The resultant mixture was stirred at ambient temperature  
25 for 1 hour. The mixture was cooled to 0°C and a 6N aqueous sodium hydroxide solution (0.8 ml) and water (2.4 ml) were added dropwise in turn. The mixture was stirred at ambient temperature for 30 minutes and then evaporated. The residue was dissolved in methylene chloride, dried over magnesium sulphate and evaporated to give 1-(2-hydroxyethyl)-5-methylimidazole (1.1 g); NMR Spectrum: (CDCl<sub>3</sub>) 2.17 (s, 3H), 3.81 (t, 2H), 3.92 (t, 2H),  
30 6.6 (s, 1H), 7.24 (s, 1H).

Tert-butyldimethylsilyl chloride (9.05 g) was added to a stirred mixture of 1-(2-hydroxyethyl)-5-methylimidazole (6.4 g), imidazole (7.5 g) and methylene chloride (30 ml) which was cooled to 0°C. The reaction mixture was stirred at ambient temperature for

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4 hours. The mixture was poured into water. The organic layer was washed with brine, dried over magnesium sulphate and evaporated to give 1-(2-~~tert~~-butyldimethylsilyloxyethyl)-5-methylimidazole (11.7 g); NMR Spectrum: (CDCl<sub>3</sub>) -0.04 (s, 6H), 0.85 (s, 6H), 2.2 (s, 3H), 3.8 (m, 2H), 3.94 (m, 2H), 6.75 (s, 1H), 7.43 (s, 1H).

5 The material so obtained was dissolved in THF (400 ml) and the solution was cooled at -60°C. n-Butyllithium (2.5M in hexane, 40 ml) was added dropwise and the mixture was stirred at -50°C for 1 hour. The mixture was cooled to -60°C and DMF (12.5 ml) was added dropwise. The resultant mixture was allowed to warm to ambient temperature and was stirred for 2 hours. Diethyl ether (500 ml) was added and the reaction mixture was poured into a  
10 saturated aqueous ammonium chloride solution. The organic layer was separated, washed with brine, dried over magnesium sulphate and evaporated. The material so obtained was purified by column chromatography on silica using increasingly polar mixtures of methylene chloride and a saturated methanolic ammonia solution as eluent. There was thus obtained 1-(2-~~tert~~-butyldimethylsilyloxyethyl)-2-formyl-5-methylimidazole (11 g); NMR Spectrum:  
15 (CDCl<sub>3</sub>) -0.1 (s, 6H), 0.79 (s, 9H), 2.32 (s, 3H), 3.91 (t, 2H), 4.4 (t, 2H), 7.07 (s, 1H), 9.71 (s, 1H).

A portion (0.79 g) of the material so obtained was dissolved in methylene chloride (24 ml) and morpholine (0.263 ml) and acetic acid (0.175 ml) were added. Sodium borohydride triacetate (0.8 g) was added portionwise and the mixture was stirred at ambient  
20 temperature for 16 hours. The mixture was evaporated and the residue was purified by column chromatography on silica using a 49:1 mixture of methylene chloride and a saturated methanolic ammonia solution as eluent. There was thus obtained 1-(2-~~tert~~-butyldimethylsilyloxyethyl)-5-methyl-2-morpholinomethylimidazole (0.5 g); NMR Spectrum: (CDCl<sub>3</sub>) 0 (s, 6H), 0.82 (s, 9H), 2.25 (s, 3H), 2.45 (m, 4H), 3.6 (s, 2H), 3.68 (m,  
25 4H), 3.85 (t, 2H), 4.1 (t, 2H), 6.7 (s, 1H).

A mixture of the material so obtained, 12N aqueous hydrochloric acid (0.26 ml) and methanol (10 ml) was stirred at ambient temperature for 5 hours. The mixture was evaporated and the residue was triturated under pentane. The resultant solid was isolated and dried under vacuum. The solid was stirred at ambient temperature for 1 hour in a mixture of methylene  
30 chloride and a saturated methanolic ammonia solution. The mixture was filtered and the filtrate was evaporated. The residue was purified by column chromatography on silica using a 19:1 mixture of methylene chloride and a saturated methanolic ammonia solution as eluent. There was thus obtained 1-(2-hydroxyethyl)-5-methyl-2-morpholinomethylimidazole (0.25 g);



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NMR Spectrum: (CDCl<sub>3</sub>) 2.2 (s, 3H), 2.6 (br s, 4H), 3.58 (s, 2H), 3.7 (m, 4H), 3.85 (t, 2H), 4.1 (t, 2H), 6.5-6.9 (br s, 1H), 6.65 (s, 1H).

**Example 7** 4-(2-chloro-5-methoxyanilino)-7-(2-hydroxy-3-pyrrolidin-1-ylpropoxy)-

5 **6-methoxyquinazoline**

A mixture of 4-(2-chloro-5-methoxyanilino)-7-(2,3-epoxypropoxy)-6-methoxyquinazoline (0.1 g), pyrrolidine (0.02 g) and chloroform (3 ml) was stirred and heated to reflux for 5 hours. The mixture was evaporated and the residue was purified by column chromatography on silica using a 4:5:1 mixture of methylene chloride, ethyl acetate  
10 and a saturated methanolic ammonia solution as eluent. There was thus obtained the title compound (0.008 g); NMR Spectrum: (DMSO-d<sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.9-2.0 (m, 2H), 2.1-2.2 (m, 2H), 3.1-3.2 (m, 2H), 3.4 (m, 2H), 3.6-3.7 (m, 2H), 3.85 (s, 3H), 4.05 (s, 3H), 4.25 (d, 2H), 4.4 (m, 1H), 7.1 (d, 1H), 7.21 (d, 1H), 7.4 (s, 1H), 7.6 (d, 1H), 8.18 (s, 1H), 8.88 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 459 and 461.

15 The 4-(2-chloro-5-methoxyanilino)-7-(2,3-epoxypropoxy)-6-methoxyquinazoline used as a starting material was prepared as follows :-

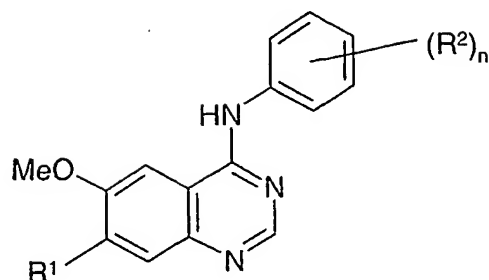
A mixture of 4-(2-chloro-5-methoxyanilino)-7-hydroxy-6-methoxyquinazoline (0.386 g), 2,3-epoxypropyl bromide (0.12 ml), potassium carbonate (0.321 g) and DMF (3 ml) was stirred and heated to 60°C for 2 hours. The mixture was cooled to ambient temperature  
20 and water (50 ml) was added. The resultant precipitate was isolated, washed with water and dried under vacuum at 55°C. The material so obtained was purified by column chromatography on silica using a 19:1 mixture of methylene chloride and methanol as eluent. There was thus obtained the required starting material (0.315 g); NMR Spectrum: (DMSO-d<sub>6</sub>) 2.78 (m, 1H), 2.9 (m, 1H), 3.45 (m, 1H), 3.8 (s, 3H), 3.95 (s, 3H), 4.0 (m, 1H), 4.53 (m, 1H),  
25 6.92 (m, 1H), 7.15 (d, 1H), 7.2 (s, 1H), 7.5 (d, 1H), 7.85 (s, 1H), 8.32 (s, 1H), 9.52 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 388 and 390.

**Example 8**

Using an analogous procedure to that described in Example 7, the appropriate  
30 7-(2,3-epoxypropoxy)quinazoline was reacted with the appropriate heterocyclic compound or amine to give the compounds described in Table III. Unless otherwise stated, each compound described in Table III was obtained as a free base and as a racemate.

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Table III



| Compound No. & Note | R <sup>1</sup>                                      | (R <sup>2</sup> ) <sub>n</sub> |
|---------------------|---|--------------------------------|
| [1]                 | 2-hydroxy-3-(4-methylpiperazin-1-yl)propoxy         | 2-chloro-5-methoxy             |
| [2]                 | 2-hydroxy-3-morpholinopropoxy                       | 2-chloro-5-methoxy             |
| [3]                 | 3-homopiperidin-1-yl-2-hydroxypropoxy               | 2-chloro-5-methoxy             |
| [4]                 | 3-[N-(2-cyanoethyl)-N-methylamino]-2-hydroxypropoxy | 2-chloro-5-methoxy             |
| [5]                 | 3-(4-acetylpiperazin-1-yl)-2-hydroxypropoxy         | 2-chloro-5-methoxy             |
| [6]                 | 3-(N-allyl-N-methylamino)-2-hydroxypropoxy          | 2-chloro-5-methoxy             |
| [7]                 | 2-hydroxy-3-(N-isopropyl-N-methylamino)propoxy      | 2-chloro-5-methoxy             |
| [8]                 | 3-azetidin-1-yl-2-hydroxypropoxy                    | 2-chloro-5-methoxy             |
| [9]                 | 3-(N-ethyl-N-isopropylamino)-2-hydroxypropoxy       | 2-chloro-5-methoxy             |
| [10]                | 2-hydroxy-3-(2-methylpyrrolidin-1-yl)propoxy        | 2-chloro-5-methoxy             |
| [11]                | 2-hydroxy-3-(1,2,5,6-tetrahydropyridin-1-yl)propoxy | 2-chloro-5-methoxy             |
| [12]                | 3-(4-cyclopropylpiperazin-1-yl)-2-hydroxypropoxy    | 2-chloro-5-methoxy             |
| [13]                | 3-(4-allylpiperazin-1-yl)-2-hydroxypropoxy          | 2-chloro-5-methoxy             |

## 5 Notes

[1] The product gave the following characterising data: NMR Spectrum: (DMSO-d<sub>6</sub>) 2.18 (s, 3H), 2.2-2.5 (m, 10H), 3.8 (s, 3H), 3.95 (s, 3H), 4.05 (d, 1H), 4.2 (d, 1H), 4.95 (br s, 1H), 6.92 (d, 1H), 7.15 (s, 1H), 7.2 (s, 1H), 7.5 (d, 1H), 7.82 (s, 1H), 8.35 (s, 1H), 9.5 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 488 and 490.

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- [2] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 3.15-3.48 (m, 4H), 3.55 (t, 2H), 3.7-3.9 (m, 2H), 3.82 (s, 3H), 4.0 (d, 2H), 4.05 (s, 3H), 4.28 (d, 2H), 4.5 (m, 1H), 7.1 (m, 1H), 7.22 (d, 1H), 7.42 (s, 1H), 7.6 (d, 1H), 8.18 (s, 1H), 8.88 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 475 and 477.
- 5 [3] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.5-1.75 (m, 4H), 1.8-2.0 (m, 4H), 3.3-3.4 (m, 3H), 3.4-3.55 (m, 3H), 3.81 (s, 3H), 4.01 (s, 3H), 4.23 (d, 2H), 4.45 (m, 1H), 7.1 (m, 1H), 7.2 (d, 1H), 7.45 (s, 1H), 7.6 (d, 1H), 8.14 (s, 1H), 8.86 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 487 and 489.
- 10 [4] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 2.3 (s, 3H), 2.45-2.75 (m, 6H), 3.8 (s, 3H), 3.98 (s, 3H), 4.0-4.1 (m, 2H), 4.2 (d, 1H), 5.0 (d, 1H), 6.92 (m, 1H), 7.15 (d, 1H), 7.2 (s, 1H), 7.5 (d, 1H), 7.82 (s, 1H), 8.32 (s, 1H), 9.5 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 472 and 474.
- 15 [5] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The reaction product was triturated under diethyl ether to which was added two equivalents of a 2M solution of hydrogen chloride in isopropanol. There was thus obtained the dihydrochloride salt of the required product which gave the following characterising data: Mass Spectrum: M+H<sup>+</sup> 516 and 518.
- 20 [6] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 2.25 (s, 3H), 2.3-2.5 (m, 2H), 3.0-3.15 (m, 2H), 3.8 (s, 3H), 3.98 (s, 3H), 4.0-4.1 (m, 2H), 4.2 (m, 1H), 4.92 (d, 1H), 5.1-5.25 (m, 2H), 5.8-5.9 (m, 1H), 6.93 (m, 1H), 7.18 (d, 1H), 7.2 (s, 1H), 7.5 (d, 1H), 7.85 (s, 1H), 8.35 (s, 1H), 9.5 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 459 and 461.
- 25 [7] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 0.95 (m, 6H), 2.25 (s, 3H), 2.4 (m, 2H), 2.8 (m, 1H), 3.8 (s, 3H), 3.9 (s, 3H), 3.9 (br s, 1H), 4.05 (m, 1H), 4.2 (m, 1H), 4.82 (br s, 1H), 6.95 (m, 1H), 7.15 (d, 1H), 7.2 (s, 1H), 7.5 (d, 1H), 7.82 (s, 1H), 8.32 (s, 1H), 9.5 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 461 and 463.
- 30 [8] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 2.0 (m, 2H), 2.45-2.55 (m, 2H), 3.2 (m, 4H), 3.8 (s, 3H), 3.8-3.9 (m, 1H), 3.98 (s, 3H), 4.03 (m, 1H), 4.12 (m, 1H), 4.92

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(d, 1H), 6.95 (m, 1H), 7.18 (d, 1H), 7.2 (s, 1H), 7.5 (d, 1H), 7.82 (s, 1H), 8.35 (s, 1H), 9.5 (s, 1H); Mass Spectrum:  $M+H^+$  445 and 447.

[9] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 0.9-1.05 (m, 9H), 2.35-2.5 (m, 2H), 2.55-2.65 (m, 2H), 2.95 (m, 1H), 3.8 (s, 3H), 3.9 (m, 1H), 3.98 (s, 3H), 4.05-4.1 (m, 1H), 4.2-4.25 (m, 1H), 4.8 (br s, 1H), 6.95 (m, 1H), 7.18 (d, 1H), 7.2 (s, 1H), 7.5 (d, 1H), 7.82 (s, 1H), 8.32 (s, 1H), 9.5 (s, 1H); Mass Spectrum:  $M+H^+$  475 and 477.

[10] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The product gave the following characterising data: Mass Spectrum:  $M+H^+$  473 and 475.

10 [11] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 2.0-2.1 (m, 2H), 2.4-2.6 (m, 4H), 2.95 (br s, 2H), 3.72 (s, 3H), 3.88 (s, 3H), 4.0 (m, 2H), 4.12 (m, 1H), 4.9 (d, 1H), 5.6 (m, 2H), 6.88 (d, 1H), 7.1 (d, 1H), 7.15 (s, 1H), 7.4 (d, 1H), 7.75 (s, 1H), 8.25 (s, 1H), 9.43 (s, 1H); Mass Spectrum:  $M+H^+$  471 and 473.

15 [12] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 0.25 (s, 2H), 0.4 (d, 2H), 1.55 (m, 1H), 2.3-2.5 (m, 10H), 3.76 (s, 3H), 3.92 (s, 3H), 4.01 (m, 2H), 4.1-4.2 (m, 1H), 4.9 (m, 1H), 6.9 (m, 1H), 7.1 (s, 1H), 7.2 (s, 1H), 7.45 (d, 1H), 7.8 (s, 1H), 8.3 (s, 1H), 9.5 (s, 1H); Mass Spectrum:  $M+H^+$  514 and 516.

20 [13] The reaction mixture was heated to 40°C for 3 hours rather than to reflux. The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 2.3-2.6 (m, 10H), 2.95 (d, 2H), 3.8 (s, 3H), 3.95 (s, 3H), 4.05 (d, 2H), 4.18 (m, 1H), 4.9 (s, 1H), 5.1-5.2 (m, 2H), 5.8 (m, 1H), 6.92 (d, 1H), 7.15 (s, 1H), 7.2 (s, 1H), 7.5 (d, 1H), 7.82 (s, 1H), 8.35 (s, 1H), 9.5 (s, 1H); Mass Spectrum:  $M+H^+$  514 and 516.

25

**Example 9 4-(2-chloro-5-methoxyanilino)-7-(2-isobutyryloxy-3-pyrrolidin-1-ylpropoxy)-6-methoxyquinazoline dihydrochloride salt**

Isobutyric acid (0.02 g) was added to a mixture of 4-(2-chloro-5-methoxyanilino)-7-(2-hydroxy-3-pyrrolidin-1-ylpropoxy)-6-methoxyquinazoline (0.1 g),  
 30 3-(3-dimethylaminopropyl)-1-ethylcarbodiimide hydrochloride (0.063 g),  
 4-dimethylaminopyridine (0.003 g) and methylene chloride (3 ml) and the mixture was stirred at ambient temperature for 16 hours. The mixture was partitioned between ethyl acetate and a 5% aqueous sodium bicarbonate solution. The organic phase was washed with water and with

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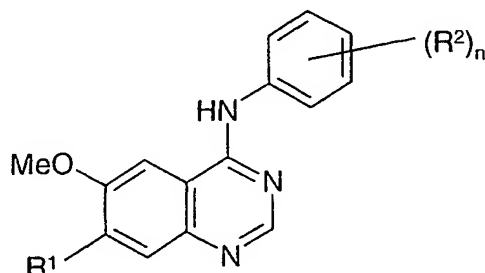
brine, dried over magnesium sulphate, and evaporated. The residue was purified by column chromatography on silica using increasingly polar solvent mixtures of methylene chloride and methanol as eluent. The material so obtained was triturated under a 6M solution of hydrogen chloride in diethyl ether. There was thus obtained the title compound (0.08 g); NMR

- 5 Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.15 (t, 6H), 1.95 (m, 2H), 2.05 (m, 2H), 2.7 (m, 1H), 3.2 (m, 2H), 3.65 (m, 4H), 3.8 (s, 3H), 4.0 (s, 3H), 4.45 (m, 2H), 5.65 (m, 1H), 7.1 (m, 1H), 7.2 (d, 1H), 7.45 (s, 1H), 7.6 (d, 1H), 8.25 (s, 1H), 8.85 (s, 1H), Mass Spectrum: [M-H]<sup>-</sup> 527 and 529.

#### 10 Example 10

Using an analogous procedure to that described in Example 9, the appropriate 7-(2-hydroxypropoxy)quinazoline was reacted with the appropriate carboxylic acid to give the compounds described in Table IV. Unless otherwise stated, each compound described in Table IV was obtained as the dihydrochloride salt.

15 **Table IV**



| Compound No. & Note | R <sup>1</sup>                                    | (R <sup>2</sup> ) <sub>n</sub> |
|---------------------|---|--------------------------------|
| [1]                 | 2-(3-methylbutyryloxy)-3-pyrrolidin-1-ylpropoxy   | 2-chloro-5-methoxy             |
| [2]                 | 2-cyclohexylcarbonyloxy-3-pyrrolidin-1-ylpropoxy  | 2-chloro-5-methoxy             |
| [3]                 | 2-cyclopentylcarbonyloxy-3-pyrrolidin-1-ylpropoxy | 2-chloro-5-methoxy             |
| [4]                 | 2-cyclobutylcarbonyloxy-3-pyrrolidin-1-ylpropoxy  | 2-chloro-5-methoxy             |

#### Notes

- 20 [1] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.9 (m, 6H), 2.0 (m, 5H), 2.35 (m, 2H), 3.2 (m, 2H), 3.65 (m, 4H), 3.8 (s, 3H), 4.0

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(s, 3H), 4.5 (m, 2H), 5.65 (m, 1H), 7.1 (m, 1H), 7.2 (d, 1H), 7.45 (s, 1H), 7.6 (d, 1H), 8.25 (s, 1H), 8.85 (s, 1H); Mass Spectrum:  $M+H^+$  541 and 543.

[2] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.2-2.1 (m, 15H), 3.2 (m, 2H), 3.6 (m, 4H), 3.7 (s, 3H), 4.0 (s, 3H), 4.45 (m, 2H), 5.6 (m, 1H), 7.1 (m, 1H), 7.2 (d, 1H), 7.5 (s, 1H), 7.6 (d, 1H), 8.25 (s, 1H), 8.85 (s, 1H); Mass Spectrum:  $M+H^+$  569 and 571.

[3] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.55-2.1 (m, 12 H), 2.9 (m, 1H), 3.2 (m, 2H), 3.65 (m, 4H), 3.8 (s, 3H), 4.0 (s, 3H), 4.4 (m, 1H), 4.5 (m, 1H), 5.6 (m, 1H), 7.1 (m, 1H), 7.2 (d, 1H), 7.45 (s, 1H), 7.6 (d, 1H), 8.2 (s, 1H), 8.85 (s, 1H); Mass Spectrum:  $M+H^+$  555 and 557.

[4] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.8-2.3 (m, 11H), 3.1-3.25 (m, 2H), 3.65 (m, 4H), 3.8 (s, 3H), 4.0 (s, 3H), 4.4 (m, 1H), 4.5 (m, 1H), 5.65 (m, 1H), 7.1 (m, 1H), 7.2 (d, 1H), 7.45 (s, 1H), 7.6 (d, 1H), 8.25 (s, 1H), 8.85 (s, 1H); Mass Spectrum:  $M+H^+$  541 and 543.

**Example 11 4-(2-bromo-5-methoxyanilino)-7-(2-hydroxy-3-pyrrolidin-1-ylpropoxy)-6-methoxyquinazoline dihydrochloride salt**

A mixture of 7-(2-acetoxy-3-pyrrolidin-1-ylpropoxy)-4-(2-bromo-5-methoxyanilino)-6-methoxyquinazoline dihydrochloride (0.123 g) and a saturated methanolic ammonia solution (3 ml) was stirred at ambient temperature for 16 hours. The mixture was evaporated and the residue was purified by column chromatography on silica (Isolute sorbent from International Sorbent Technology Ltd, ref 9470-0100) using a 19:1 mixture of methylene chloride and methanol as eluent. The material so obtained was dissolved in methylene chloride and a 6M solution of hydrogen chloride in isopropanol (0.3 ml) was added. The mixture was diluted with diethyl ether (10 ml) and the resultant solid was collected, washed with diethyl ether and dried under vacuum. There was thus obtained the title compound (0.115 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.9 (m, 2H), 2.05 (m, 2H), 3.15 (m, 2H), 3.4 (m, 2H), 3.65 (m, 2H), 3.8 (s, 3H), 4.05 (s, 3H), 4.25 (d, 2H), 4.4 (m, 1H), 7.05 (m, 1H), 7.2 (d, 1H), 7.45 (s, 1H), 7.7 (d, 1H), 8.25 (s, 1H), 8.85 (s, 1H); Mass Spectrum:  $M+H^+$  503 and 505.

The 7-(2-acetoxy-3-pyrrolidin-1-ylpropoxy)-4-(2-bromo-5-methoxyanilino)-6-methoxyquinazoline dihydrochloride used as a starting material was prepared as follows :-

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7-(2,3-Epoxypropoxy)-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one was reacted with pyrrolidine using an analogous procedure to that described in the second paragraph of the portion of Note [6] in Example 4 above that is concerned with the preparation of starting materials. There was thus obtained 7-(2-hydroxy-3-pyrrolidin-1-ylpropoxy)-

5 6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one.

The material so obtained was taken through an analogous sequence of reactions to those described in the third to fifth paragraphs of the portion of Note [6] in Example 4 above that is concerned with the preparation of starting materials. There was thus obtained

7-(2-acetoxy-3-pyrrolidin-1-ylpropoxy)-4-chloro-6-methoxyquinazoline; NMR Spectrum:  
10 (CDCl<sub>3</sub> and CD<sub>3</sub>CO<sub>2</sub>D) 2.05 (s, 4H), 2.15 (s, 3H), 3.45 (br s, 4H), 3.65 (m, 2H), 4.05 (s, 3H), 4.4 (d, 2H), 5.65 (m, 1H), 7.4 (s, 1H), 7.55 (s, 1H), 8.9 (s, 1H).

The material so obtained was reacted with 2-bromo-5-methoxyaniline using an analogous procedure to that described in Example 3. There was thus obtained 7-(2-acetoxy-3-pyrrolidin-1-ylpropoxy)-4-(2-bromo-5-methoxyanilino)-6-methoxyquinazoline

15 dihydrochloride salt; Mass Spectrum: M+H<sup>+</sup> 545 and 547.

**Example 12 4-(2-bromo-5-methoxyanilino)-7-[2-hydroxy-3-(N-isopropyl-N-methylamino)propoxy]-6-methoxyquinazoline dihydrochloride salt**

Using an analogous procedure to that described in Example 11, 7-[2-acetoxy-  
20 3-(N-isopropyl-N-methylamino)propoxy]-4-(2-bromo-5-methoxyanilino)-6-methoxyquinazoline dihydrochloride was reacted with a saturated methanolic ammonia solution to give the title compound; Mass Spectrum: M+H<sup>+</sup> 505 and 507.

The 7-[2-acetoxy-3-(N-isopropyl-N-methylamino)propoxy]-4-(2-bromo-5-methoxyanilino)-6-methoxyquinazoline dihydrochloride used as a starting material was  
25 prepared as follows :-

7-(2,3-Epoxypropoxy)-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one was reacted with N-isopropyl-N-methylamine using an analogous procedure to that described in the second paragraph of the portion of Note [6] in Example 4 above that is concerned with the preparation of starting materials. There was thus obtained 7-[2-hydroxy-3-(N-isopropyl-N-methylamino)propoxy]-6-methoxy-3-pivaloyloxymethyl-3,4-dihydroquinazolin-4-one.  
30

The material so obtained was taken through an analogous sequence of reactions to those described in the third to fifth paragraphs of the portion of Note [6] in Example 4 above that is concerned with the preparation of starting materials. There was thus obtained

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7-[2-acetoxy-3-(N-isopropyl-N-methyamino)propoxy]-4-chloro-6-methoxyquinazoline; NMR Spectrum: (CDCl<sub>3</sub>) 1.0 (d, 6H), 2.1 (s, 3H), 2.3 (s, 3H), 2.6 (m, 1H), 2.75 (m, 1H), 2.85 (m, 1H), 4.05 (s, 3H), 4.35 (m, 1H), 4.45 (m, 1H), 5.35 (m, 1H), 7.25 (s, 1H), 7.4 (s, 1H), 8.85 (s, 1H).

5 The material so obtained was reacted with 2-bromo-5-methoxyaniline using an analogous procedure to that described in Example 3. There was thus obtained 7-[2-acetoxy-3-(N-isopropyl-N-methyamino)propoxy]-4-(2-bromo-5-methoxyanilino)-6-methoxyquinazoline dihydrochloride salt; Mass Spectrum: M+H<sup>+</sup> 547 and 549.

10 **Example 13** 4-(2-chloro-5-methoxyanilino)-7-[(2R)-(2-hydroxy-3-morpholinopropoxy)]-6-methoxyquinazoline dihydrochloride salt

A mixture of 4-(2-chloro-5-methoxyanilino)-7-[(2R)-2,3-epoxypropoxy]-6-methoxyquinazoline (0.1 g), morpholine (2.5 equivalents), chloroform (2.5 ml) and ethanol (2.5 ml) was stirred and heated to 40°C for 8 hours. The solvent was evaporated. Methylene chloride (5 ml) and a polystyrene isocyanate resin (0.3 g; loading: 1 mmol/g; prepared according to the procedure disclosed in J. Amer. Chem. Soc., 1997, 119, 4882) was added and the mixture was stirred at ambient temperature for 1.5 hours. The mixture was filtered and the filtrate was evaporated. The crude product so obtained was purified by column chromatography on silica using increasingly polar mixtures of methylene chloride and a saturated methanolic ammonia solution as eluent. There was thus obtained the title compound; NMR Spectrum: (DMSO-d<sub>6</sub>) 2.4-2.6 (m, 6H), 3.6 (m, 4H), 3.8 (s, 3H), 3.95 (s, 3H), 4.05 (d, 2H), 4.2 (m, 1H), 4.95 (d, 1H), 6.9 (m, 1H), 7.15 (d, 1H), 7.2 (s, 1H), 7.5 (d, 1H), 7.8 (s, 1H), 8.3 (s, 1H), 9.5 (s, 1H); Mass Spectrum: M+H<sup>+</sup> 475 and 477.

The 4-(2-chloro-5-methoxyanilino)-7-[(2R)-2,3-epoxypropoxy]-6-methoxyquinazoline used as a starting material was prepared as follows :-

(2R)-(-)-Glycidyl tosylate (3.6 g) was added to a mixture of 4-(2-chloro-5-methoxyanilino)-7-hydroxy-6-methoxyquinazoline (4.8 g), potassium carbonate (8.5 g) and DMF (60 ml) and the mixture was stirred and heated to 40°C for 4 hours. The resultant mixture was filtered and the filtrate was evaporated. The residue was partitioned between methylene chloride and water. The organic phase was washed in turn with a 5% aqueous ammonium hydroxide solution, with water and with brine, dried over magnesium sulphate and evaporated. The material so obtained was washed with a 4:1 mixture of petroleum ether and diethyl ether and dried under vacuum. There was thus obtained 4-(2-chloro-



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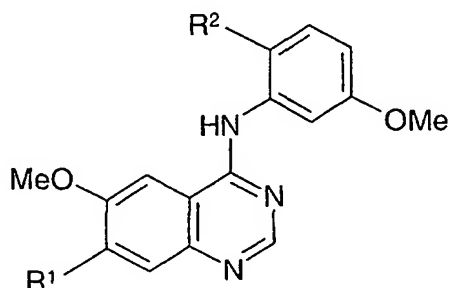
5-methoxyanilino)-7-[(2R)-2,3-epoxypropoxy]-6-methoxyquinazoline as a solid (3.1 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 2.8 (m, 1H), 2.9 (m, 1H), 3.45 (m, 1H), 3.8 (s, 3H), 3.95 (s, 3H), 4.0 (m, 1H), 4.55 (m, 1H), 6.9 (m, 1H), 7.15 (d, 1H), 7.2 (s, 1H), 7.55 (d, 1H), 7.85 (s, 1H), 8.3 (s, 1H), 9.5 (br s, 1H).

5

**Example 14**

Using an analogous procedure to that described in Example 13, the appropriate 7-(2,3-epoxypropoxy)quinazoline was reacted with the appropriate heterocyclic compound or amine to give the compounds described in Table V. In each case, the reaction product was dissolved in a 9:1 mixture of methylene chloride and methanol and a 2.2M hydrogen chloride solution in diethyl ether was added. Each precipitate was isolated and dried under vacuum to give the desired products as dihydrochloride salts.

Table V



15

| Compound No. & Note | R <sup>1</sup>                                       | R <sup>2</sup> |
|---------------------|--|----------------|
| [1]                 | (2R)-3-(N-allyl-N-cyclopentylamino)-2-hydroxypropoxy | chloro         |
| [2]                 | (2R)-3-(N-allyl-N-methylamino)-2-hydroxypropoxy      | chloro         |
| [3]                 | (2R)-2-hydroxy-3-(N-isobutyl-N-methylamino)propoxy   | bromo          |
| [4]                 | (2R)-2-hydroxy-3-piperidinopropoxy                   | bromo          |

Notes

[1] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.5-1.65 (m, 2H), 1.68-1.96 (m, 4H), 2.0-2.17 (m, 2H), 3.25-3.43 (m, 2H), 3.75 (m, 1H), 3.82 (s, 3H), 3.92 (d, 2H), 3.95 (m, 1H), 4.0 (s, 3H), 4.2-4.35 (m, 2H), 4.43-4.58 (m, 1H), 5.53-5.73 (m, 2H), 6.02-6.18 (m, 1H), 7.1 (m, 1H), 7.2 (d, 1H), 7.45 (s, 1H), 7.6 (d, 1H), 8.25 (s, 1H), 8.85 (s, 1H); Mass Spectrum: M-H 511 and 513.

20

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- [2] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 2.85 (br s, 3H), 3.19-3.44 (m, 2H), 3.81 (s, 3H), 3.77-3.97 (m, 2H), 4.03 (s, 3H), 4.2-4.3 (m, 2H), 4.44-4.56 (m, 1H), 5.51-5.64 (m, 2H), 5.95-6.08 (m, 1H), 7.08 (m, 1H), 7.2 (d, 1H), 7.5 (br s, 1H), 7.57 (d, 1H), 8.3 (br s, 1H), 8.84 (s, 1H); Mass Spectrum: M-H<sup>+</sup> 457 and 459.
- [3] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 0.93-1.1 (m, 6H), 2.06-2.22 (m, 1H), 2.9 (s, 3H), 2.9-3.03 (m, 1H), 3.09-3.48 (m, 3H), 3.81 (s, 3H), 4.03 (s, 3H), 4.2-4.31 (m, 2H), 4.46-4.58 (m, 1H), 7.02 (m, 1H), 1.19 (d, 1H), 7.49 (s, 1H), 7.72 (d, 1H), 8.26 (s, 1H), 8.84 (s, 1H); Mass Spectrum: M-H<sup>+</sup> 517 and 519.
- 10 The 4-(2-bromo-5-methoxyanilino)-7-[(2R)-2,3-epoxypropoxy]-6-methoxyquinazoline used as a starting material was prepared as follows :-
- A 1M solution in THF of the sodium salt of 1,1,1,3,3,3-hexamethyldisilazane (82.4 ml) was added dropwise to a mixture of 2-bromo-5-methoxyaniline (16.7 g) and DMF (200 ml) and the reaction mixture was stirred at ambient temperature for a further 30 minutes.
- 15 A solution of 7-benzyloxy-4-chloro-6-methoxyquinazoline (11.8 g) in DMF (250 ml) was added and the reaction mixture was at ambient temperature for 20 minutes. The mixture was concentrated by evaporation of about half of the DMF. The residue was partitioned between methylene chloride and water. The organic phase was dried over magnesium sulphate and evaporated. The resultant solid was washed with a 1:1 mixture of petroleum ether and diethyl
- 20 ether and dried overnight under vacuum. There was thus obtained 4-(2-bromo-5-methoxyanilino)-7-benzyloxy-6-methoxyquinazoline (13.1 g); NMR Spectrum: (DMSO<sub>d</sub><sub>6</sub>) 3.8 (s, 3H), 3.95 (s, 3H), 5.3 (s, 2H), 6.85 (m, 1H), 7.15 (d, 1H), 7.3 (s, 1H), 7.35-7.55 (m, 5H), 7.6 (d, 1H), 7.85 (s, 1H), 8.3 (s, 1H), 9.5 (br s, 1H).
- A mixture of the material so obtained and trifluoroacetic acid (130 ml) was stirred and
- 25 heated to reflux for 5 hours. The mixture was evaporated, water was added and the mixture was neutralised by the addition of a saturated aqueous sodium bicarbonate solution. The resultant solid was dried under vacuum to give 4-(2-bromo-5-methoxyanilino)-7-hydroxy-6-methoxyquinazoline (11 g); Mass Spectrum: M-H<sup>+</sup> 374 and 376.
- (2R)-(-)-Glycidyl tosylate (3.6 g) was added to a mixture of 4-(2-bromo-
- 30 5-methoxyanilino)-7-hydroxy-6-methoxyquinazoline (5 g), potassium carbonate (7.3 g) and DMF (60 ml) and the mixture was stirred and heated to 40°C for 4 hours. The resultant mixture was filtered and the filtrate was evaporated. The residue was partitioned between methylene chloride and water. The organic phase was washed in turn with a 5% aqueous

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ammonium hydroxide solution, with water and with brine, dried over magnesium sulphate and evaporated. The material so obtained was washed with a 4:1 mixture of petroleum ether and diethyl ether and dried under vacuum. There was thus obtained 4-(2-bromo-

5-methoxyanilino)-7-[(2R)-2,3-epoxypropoxy]-6-methoxyquinazoline as a solid (3.6 g); NMR Spectrum: (DMSO<sub>d6</sub>) 2.8 (m, 1H), 2.9 (m, 1H), 3.45 (m, 1H), 3.8 (s, 3H), 3.95 (s, 3H), 4.05 (m, 1H), 4.55 (m, 1H), 6.9 (m, 1H), 7.15 (d, 1H), 7.2 (s, 1H), 7.6 (d, 1H), 7.85 (s, 1H), 8.3 (s, 1H), 9.5 (br s, 1H); Mass Spectrum: M+H<sup>+</sup> 432 and 434.

[4] The product gave the following characterising data: NMR Spectrum: (DMSO<sub>d6</sub> and CF<sub>3</sub>CO<sub>2</sub>D) 1.33-1.52 (m, 1H), 1.64-1.97 (m, 5H), 2.93-3.14 (m, 2H), 3.2-3.4 (m, 2H), 3.47-3.64 (m, 2H), 3.82 (s, 3H), 4.04 (s, 3H), 4.26 (s, 2H), 4.46-4.59 (m, 1H), 7.03 (m, 1H), 7.21 (d, 1H), 7.46 (s, 1H), 7.73 (d, 1H), 8.21 (s, 1H), 8.56 (s, 1H); Mass Spectrum: M-H<sup>+</sup> 515 and 517.

### Example 15

#### 15 **Pharmaceutical compositions**

The following illustrate representative pharmaceutical dosage forms of the invention as defined herein (the active ingredient being termed "Compound X"), for therapeutic or prophylactic use in humans:

|    |     |  |           |
|----|-----|--|-----------|
| 20 | (a) | Tablet I                                 | mg/tablet |
|    |     | Compound X.....                          | 100       |
|    |     | Lactose Ph.Eur.....                      | 182.75    |
|    |     | Croscarmellose sodium.....               | 12.0      |
|    |     | Maize starch paste (5% w/v paste).....   | 2.25      |
| 25 |     | Magnesium stearate.....                  | 3.0       |
|    | (b) | Tablet II                                | mg/tablet |
|    |     | Compound X.....                          | 50        |
|    |     | Lactose Ph.Eur.....                      | 223.75    |
| 30 |     | Croscarmellose sodium.....               | 6.0       |
|    |     | Maize starch.....                        | 15.0      |
|    |     | Polyvinylpyrrolidone (5% w/v paste)..... | 2.25      |
|    |     | Magnesium stearate.....                  | 3.0       |

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|    |     |  |                           |
|----|-----|--|---------------------------|
| 5  | (c) | Tablet III                                   | mg/tablet                 |
|    |     | Compound X.....                              | 1.0                       |
|    |     | Lactose Ph.Eur.....                          | 93.25                     |
|    |     | Croscarmellose sodium.....                   | 4.0                       |
|    |     | Maize starch paste (5% w/v paste).....       | 0.75                      |
|    |     | Magnesium stearate.....                      | 1.0                       |
| 10 | (d) | Capsule                                      | mg/capsule                |
|    |     | Compound X.....                              | 10                        |
|    |     | Lactose Ph.Eur.....                          | 488.5                     |
|    |     | Magnesium.....                               | 1.5                       |
| 15 | (e) | Injection I                                  | (50 mg/ml)                |
|    |     | Compound X.....                              | 5.0% w/v                  |
|    |     | 1M Sodium hydroxide solution.....            | 15.0% v/v                 |
|    |     | 0.1M Hydrochloric acid (to adjust pH to 7.6) |                           |
|    |     | Polyethylene glycol 400.....                 | 4.5% w/v                  |
|    |     | Water for injection to 100%                  |                           |
| 20 | (f) | Injection II                                 | (10 mg/ml)                |
|    |     | Compound X.....                              | 1.0% w/v                  |
|    |     | Sodium phosphate BP.....                     | 3.6% w/v                  |
|    |     | 0.1M Sodium hydroxide solution.....          | 15.0% v/v                 |
|    |     | Water for injection to 100%                  |                           |
| 25 |     |  |                           |
| 30 | (g) | Injection III                                | (1mg/ml, buffered to pH6) |
|    |     | Compound X.....                              | 0.1% w/v                  |
|    |     | Sodium phosphate BP.....                     | 2.26% w/v                 |
|    |     | Citric acid.....                             | 0.38% w/v                 |
|    |     | Polyethylene glycol 400.....                 | 3.5% w/v                  |
|    |     | Water for injection to 100%                  |                           |

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|    |     |                                    |        |
|----|-----|------------------------------------|--------|
| 5  | (h) | Aerosol I                          | mg/ml  |
|    |     | Compound X.....                    | 10.0   |
|    |     | Sorbitan trioleate.....            | 13.5   |
|    |     | Trichlorofluoromethane.....        | 910.0  |
|    |     | Dichlorodifluoromethane.....       | 490.0  |
| 10 | (i) | Aerosol II                         | mg/ml  |
|    |     | Compound X.....                    | 0.2    |
|    |     | Sorbitan trioleate.....            | 0.27   |
|    |     | Trichlorofluoromethane.....        | 70.0   |
|    |     | Dichlorodifluoromethane.....       | 280.0  |
|    |     | Dichlorotetrafluoroethane.....     | 1094.0 |
| 15 | (j) | Aerosol III                        | mg/ml  |
|    |     | Compound X.....                    | 2.5    |
|    |     | Sorbitan trioleate.....            | 3.38   |
|    |     | Trichlorofluoromethane.....        | 67.5   |
|    |     | Dichlorodifluoromethane.....       | 1086.0 |
|    |     | Dichlorotetrafluoroethane.....     | 191.6  |
| 20 | (k) | Aerosol IV                         | mg/ml  |
|    |     | Compound X.....                    | 2.5    |
|    |     | Soya lecithin.....                 | 2.7    |
|    |     | Trichlorofluoromethane.....        | 67.5   |
|    |     | Dichlorodifluoromethane.....       | 1086.0 |
|    |     | Dichlorotetrafluoroethane.....     | 191.6  |
| 30 | (l) | Ointment                           | ml     |
|    |     | Compound X.....                    | 40 mg  |
|    |     | Ethanol.....                       | 300 µl |
|    |     | Water.....                         | 300 µl |
|    |     | 1-Dodecylazacycloheptan-2-one..... | 50 µl  |

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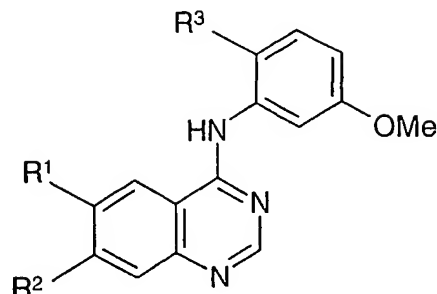
Propylene glycol..... to 1 ml

Note

The above formulations may be obtained by conventional procedures well known in the pharmaceutical art. The tablets (a)-(c) may be enteric coated by conventional means, for example to provide a coating of cellulose acetate phthalate. The aerosol formulations (h)-(k) may be used in conjunction with standard, metered dose aerosol dispensers, and the suspending agents sorbitan trioleate and soya lecithin may be replaced by an alternative suspending agent such as sorbitan monooleate, sorbitan sesquioleate, polysorbate 80, polyglycerol oleate or oleic acid.

CLAIMS

1. A quinazoline derivative of the Formula I

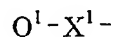


I

- 5 wherein :-

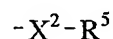
$R^3$  is chloro, bromo or iodo;

$R^1$  is hydrogen or (1-6C)alkoxy and  $R^2$  is a group of the formula :



- 10 wherein  $X^1$  is selected from O, S, SO, SO<sub>2</sub>, N(R<sup>4</sup>), CO, CH(OR<sup>4</sup>), CON(R<sup>4</sup>), N(R<sup>4</sup>)CO, SO<sub>2</sub>N(R<sup>4</sup>), N(R<sup>4</sup>)SO<sub>2</sub>, OC(R<sup>4</sup>)<sub>2</sub>, SC(R<sup>4</sup>)<sub>2</sub> and N(R<sup>4</sup>)C(R<sup>4</sup>)<sub>2</sub>, wherein R<sup>4</sup> is hydrogen or (1-6C)alkyl, and Q<sup>1</sup> is heteroaryl, heteroaryl-(1-6C)alkyl, heteroaryloxy-(1-6C)alkyl, heterocyclyl, heterocyclyl-(1-6C)alkyl or heterocyclyloxy-(1-6C)alkyl, provided that, when X<sup>1</sup> is O, any compound wherein Q<sup>1</sup> is a piperidino-(2-4C)alkyl, morpholino-(2-4C)alkyl or
- 15 piperazin-1-yl-(2-4C)alkyl group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is excluded, and provided that, when X<sup>1</sup> is O, any compound wherein Q<sup>1</sup> is a 4-(1-4C)alkylpiperazin-1-yl-(2-4C)alkyl group that bears no further substituent on the 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is also excluded,

- 20 or  $R^2$  is a group of the formula :



- wherein X<sup>2</sup> is selected from O and N(R<sup>6</sup>), wherein R<sup>6</sup> is hydrogen or (1-6C)alkyl, and R<sup>5</sup> is hydroxy-(3-6C)alkyl, (1-6C)alkoxy-(3-6C)alkyl, amino-(3-6C)alkyl, (1-6C)alkylamino-(3-6C)alkyl, di-[(1-6C)alkyl]amino-(3-6C)alkyl, (2-6C)alkanoylamino-(3-6C)alkyl or
- 25 (1-6C)alkoxycarbonylamino-(3-6C)alkyl, provided that the central CH<sub>2</sub> group within any (CH<sub>2</sub>)<sub>3</sub> group within the (3-6C)alkylene group in a R<sup>5</sup> group bears a substituent selected from hydroxy, amino, (1-6C)alkoxy, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (2-6C)alkanoyloxy, (2-6C)alkanoylamino and N-(1-6C)alkyl-(2-6C)alkanoylamino,

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and wherein adjacent carbon atoms in any (2-6C)alkylene chain within a  $R^2$  substituent are optionally separated by the insertion into the chain of a group selected from O, S, SO, SO<sub>2</sub>, N(R<sup>7</sup>), CO, CH(OR<sup>7</sup>), CON(R<sup>7</sup>), N(R<sup>7</sup>)CO, SO<sub>2</sub>N(R<sup>7</sup>), N(R<sup>7</sup>)SO<sub>2</sub>, CH=CH and C≡C wherein R<sup>7</sup> is hydrogen or (1-6C)alkyl, or, when the inserted group is N(R<sup>7</sup>), R<sup>7</sup> may also be

5 (2-6C)alkanoyl,

and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a  $R^2$  substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group one or more halogeno or (1-6C)alkyl substituents or a substituent selected from hydroxy, cyano, amino, carboxy, carbamoyl, (2-8C)alkenyl, (2-8C)alkynyl, (1-6C)alkoxy, (1-6C)alkylthio, (1-6C)alkylsulphinyl, (1-6C)alkylsulphonyl, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (1-6C)alkoxycarbonyl, N-(1-6C)alkylcarbamoyl, N,N-di-[(1-6C)alkyl]carbamoyl, (2-6C)alkanoyl, (2-6C)alkanoyloxy, (2-6C)alkanoylamino, N-(1-6C)alkyl-(2-6C)alkanoylamino, N-(1-6C)alkylsulphamoyl, N,N-di-[(1-6C)alkyl]sulphamoyl, (1-6C)alkanesulphonylamino and N-(1-6C)alkyl-(1-6C)alkanesulphonylamino, or from a group of the formula :

15  $-X^4-Q^2$

wherein X<sup>4</sup> is a direct bond or is selected from O, S, SO, SO<sub>2</sub>, N(R<sup>8</sup>), CO, CH(OR<sup>8</sup>), CON(R<sup>8</sup>), N(R<sup>8</sup>)CO, SO<sub>2</sub>N(R<sup>8</sup>), N(R<sup>8</sup>)SO<sub>2</sub>, C(R<sup>8</sup>)<sub>2</sub>O, C(R<sup>8</sup>)<sub>2</sub>S and N(R<sup>8</sup>)C(R<sup>8</sup>)<sub>2</sub>, wherein R<sup>8</sup> is hydrogen or (1-6C)alkyl, and Q<sup>2</sup> is aryl, aryl-(1-6C)alkyl, (3-7C)cycloalkyl, (3-7C)cycloalkyl-(1-6C)alkyl, (3-7C)cycloalkenyl, (3-7C)cycloalkenyl-(1-6C)alkyl, heteroaryl, heteroaryl-(1-6C)alkyl, heterocyclyl or heterocyclyl-(1-6C)alkyl,

and wherein any aryl, heteroaryl or heterocyclyl group within a substituent on  $R^2$  optionally bears 1, 2 or 3 substituents, which may be the same or different, selected from halogeno, trifluoromethyl, cyano, nitro, hydroxy, amino, carboxy, carbamoyl, (1-6C)alkyl, (2-8C)alkenyl, (2-8C)alkynyl, (1-6C)alkoxy, (2-6C)alkenyloxy, (2-6C)alkynyloxy, (1-6C)alkylthio, (1-6C)alkylsulphinyl, (1-6C)alkylsulphonyl, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (1-6C)alkoxycarbonyl, N-(1-6C)alkylcarbamoyl, N,N-di-[(1-6C)alkyl]carbamoyl, (2-6C)alkanoyl, (2-6C)alkanoyloxy, (2-6C)alkanoylamino, N-(1-6C)alkyl-(2-6C)alkanoylamino, N-(1-6C)alkylsulphamoyl, N,N-di-[(1-6C)alkyl]sulphamoyl, (1-6C)alkanesulphonylamino and N-(1-6C)alkyl-(1-6C)alkanesulphonylamino, or from a group of the formula :

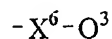
30  $-X^5-R^9$



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wherein  $X^5$  is a direct bond or is selected from O and  $N(R^{10})$ , wherein  $R^{10}$  is hydrogen or (1-6C)alkyl, and  $R^9$  is halogeno-(1-6C)alkyl, hydroxy-(1-6C)alkyl, (1-6C)alkoxy-(1-6C)alkyl, cyano-(1-6C)alkyl, amino-(1-6C)alkyl, (1-6C)alkylamino-(1-6C)alkyl, di-[(1-6C)alkyl]amino-(1-6C)alkyl, (2-6C)alkanoylamino-(1-6C)alkyl or (1-6C)alkoxycarbonylamino-(1-6C)alkyl,

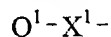
5 or from a group of the formula :



wherein  $X^6$  is a direct bond or is selected from O, CO and  $N(R^{11})$ , wherein  $R^{11}$  is hydrogen or (1-6C)alkyl, and  $Q^3$  is aryl, aryl-(1-6C)alkyl, heteroaryl, heteroaryl-(1-6C)alkyl, heterocyclyl or heterocyclyl-(1-6C)alkyl which optionally bears 1 or 2 substituents, which may be the same  
10 or different, selected from halogeno, (1-6C)alkyl, (2-8C)alkenyl, (2-8C)alkynyl and (1-6C)alkoxy,

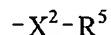
and wherein any heterocyclyl group within a substituent on  $R^2$  optionally bears 1 or 2 oxo or thioxo substituents;

15 or wherein  $R^2$  is hydrogen or (1-6C)alkoxy and  $R^1$  is a group of the formula :



wherein  $X^1$  is selected from O, S, SO,  $SO_2$ ,  $N(R^4)$ , CO,  $CH(OR^4)$ ,  $CON(R^4)$ ,  $N(R^4)CO$ ,  $SO_2N(R^4)$ ,  $N(R^4)SO_2$ ,  $OC(R^4)_2$ ,  $SC(R^4)_2$  and  $N(R^4)C(R^4)_2$ , wherein  $R^4$  is hydrogen or (1-6C)alkyl, and  $Q^1$  is heterocyclyl, heterocyclyl-(1-6C)alkyl or heterocyclyloxy-(1-6C)alkyl,  
20 provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a piperidino-(2-4C)alkyl, morpholino-(2-4C)alkyl or piperazin-1-yl-(2-4C)alkyl group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is excluded, and provided that, when  $X^1$  is O, any compound wherein  $Q^1$  is a 4-(1-4C)alkylpiperazin-1-yl-(2-4C)alkyl group that bears no further substituent on the  
25 4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the (2-4C)alkylene group is also excluded,

or  $R^1$  is a group of the formula :



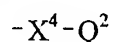
wherein  $X^2$  is selected from O and  $N(R^6)$ , wherein  $R^6$  is hydrogen or (1-6C)alkyl, and  $R^5$  is  
30 hydroxy-(3-6C)alkyl, (1-6C)alkoxy-(3-6C)alkyl, amino-(3-6C)alkyl, (1-6C)alkylamino-(3-6C)alkyl, di-[(1-6C)alkyl]amino-(3-6C)alkyl, (2-6C)alkanoylamino-(3-6C)alkyl or (1-6C)alkoxycarbonylamino-(3-6C)alkyl, provided that the central  $CH_2$  group within any  $(CH_2)_3$  group within the (3-6C)alkylene group in a  $R^5$  group bears a substituent selected from

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hydroxy, amino, (1-6C)alkoxy, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (2-6C)alkanoyloxy, (2-6C)alkanoylamino and N-(1-6C)alkyl-(2-6C)alkanoylamino,

and wherein adjacent carbon atoms in any (2-6C)alkylene chain within a  $R^1$  substituent are optionally separated by the insertion into the chain of a group selected from O, S, SO, SO<sub>2</sub>,  
 5 N( $R^7$ ), CO, CH(OR<sup>7</sup>), CON( $R^7$ ), N( $R^7$ )CO, SO<sub>2</sub>N( $R^7$ ), N( $R^7$ )SO<sub>2</sub>, CH=CH and C≡C wherein  $R^7$  is hydrogen or (1-6C)alkyl, or, when the inserted group is N( $R^7$ ),  $R^7$  may also be (2-6C)alkanoyl,

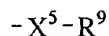
and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a  $R^1$  substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group one or more halogeno or (1-6C)alkyl substituents or a substituent  
 10 selected from hydroxy, cyano, amino, carboxy, carbamoyl, (2-8C)alkenyl, (2-8C)alkynyl, (1-6C)alkoxy, (1-6C)alkylthio, (1-6C)alkylsulphinyl, (1-6C)alkylsulphonyl, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (1-6C)alkoxycarbonyl, N-(1-6C)alkylcarbamoyl, N,N-di-[(1-6C)alkyl]carbamoyl, (2-6C)alkanoyl, (2-6C)alkanoyloxy, (2-6C)alkanoylamino, N-(1-6C)alkyl-(2-6C)alkanoylamino, N-(1-6C)alkylsulphamoyl,  
 15 N,N-di-[(1-6C)alkyl]sulphamoyl, (1-6C)alkanesulphonylamino and N-(1-6C)alkyl-(1-6C)alkanesulphonylamino, or from a group of the formula :



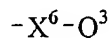
wherein  $X^4$  is a direct bond or is selected from O, S, SO, SO<sub>2</sub>, N( $R^8$ ), CO, CH(OR<sup>8</sup>), CON( $R^8$ ), N( $R^8$ )CO, SO<sub>2</sub>N( $R^8$ ), N( $R^8$ )SO<sub>2</sub>, C( $R^8$ )<sub>2</sub>O, C( $R^8$ )<sub>2</sub>S and N( $R^8$ )C( $R^8$ )<sub>2</sub>, wherein  $R^8$  is  
 20 hydrogen or (1-6C)alkyl, and  $Q^2$  is aryl, aryl-(1-6C)alkyl, (3-7C)cycloalkyl, (3-7C)cycloalkyl-(1-6C)alkyl, (3-7C)cycloalkenyl, (3-7C)cycloalkenyl-(1-6C)alkyl, heteroaryl, heteroaryl-(1-6C)alkyl, heterocyclyl or heterocyclyl-(1-6C)alkyl,

and wherein any aryl, heteroaryl or heterocyclyl group within a substituent on  $R^1$  optionally bears 1, 2 or 3 substituents, which may be the same or different, selected from  
 25 halogeno, trifluoromethyl, cyano, nitro, hydroxy, amino, carboxy, carbamoyl, (1-6C)alkyl, (2-8C)alkenyl, (2-8C)alkynyl, (1-6C)alkoxy, (2-6C)alkenyloxy, (2-6C)alkynyloxy, (1-6C)alkylthio, (1-6C)alkylsulphinyl, (1-6C)alkylsulphonyl, (1-6C)alkylamino, di-[(1-6C)alkyl]amino, (1-6C)alkoxycarbonyl, N-(1-6C)alkylcarbamoyl, N,N-di-[(1-6C)alkyl]carbamoyl, (2-6C)alkanoyl, (2-6C)alkanoyloxy, (2-6C)alkanoylamino,  
 30 N-(1-6C)alkyl-(2-6C)alkanoylamino, N-(1-6C)alkylsulphamoyl, N,N-di-[(1-6C)alkyl]sulphamoyl, (1-6C)alkanesulphonylamino and N-(1-6C)alkyl-(1-6C)alkanesulphonylamino, or from a group of the formula :

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wherein  $X^5$  is a direct bond or is selected from O and N( $R^{10}$ ), wherein  $R^{10}$  is hydrogen or (1-6C)alkyl, and  $R^9$  is halogeno-(1-6C)alkyl, hydroxy-(1-6C)alkyl, (1-6C)alkoxy-(1-6C)alkyl, cyano-(1-6C)alkyl, amino-(1-6C)alkyl, (1-6C)alkylamino-(1-6C)alkyl, di-[(1-6C)alkyl]amino-  
 5 (1-6C)alkyl, (2-6C)alkanoylamino-(1-6C)alkyl or (1-6C)alkoxycarbonylamino-(1-6C)alkyl, or from a group of the formula :



wherein  $X^6$  is a direct bond or is selected from O, CO and N( $R^{11}$ ), wherein  $R^{11}$  is hydrogen or (1-6C)alkyl, and  $Q^3$  is aryl, aryl-(1-6C)alkyl, heteroaryl, heteroaryl-(1-6C)alkyl, heterocyclyl  
 10 or heterocyclyl-(1-6C)alkyl which optionally bears 1 or 2 substituents, which may be the same or different, selected from halogeno, (1-6C)alkyl, (2-8C)alkenyl, (2-8C)alkynyl and (1-6C)alkoxy,

and wherein any heterocyclyl group within a substituent on  $R^1$  optionally bears 1 or 2 oxo or thioxo substituents;

15 or a pharmaceutically-acceptable salt thereof.

2. A quinazoline derivative of the Formula I according to claim 1 wherein :

$R^1$  is methoxy and  $R^2$  is 2-imidazol-1-ylethoxy, 3-(1,2,3-triazol-1-yl)propoxy, 4-pyridylmethoxy, 2-pyrid-4-yloxyethoxy, 3-pyrid-2-yloxypropoxy, 3-pyrid-4-yloxypropoxy,  
 20 2-pyrrolidin-1-ylethoxy, 3-pyrrolidin-1-ylpropoxy, pyrrolidin-3-yloxy, pyrrolidin-2-ylmethoxy, 2-pyrrolidin-2-ylethoxy, 3-pyrrolidin-2-ylpropoxy, 2-morpholinoethoxy, 3-morpholinopropoxy, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)ethoxy, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-piperidinoethoxy, 3-piperidinopropoxy, piperidin-3-yloxy, piperidin-4-yloxy, piperidin-3-ylmethoxy,  
 25 2-piperidin-3-ylethoxy, piperidin-4-ylmethoxy, 2-piperidin-4-ylethoxy, 2-homopiperidin-1-ylethoxy, 3-homopiperidin-1-ylpropoxy, 2-piperazin-1-ylethoxy, 3-piperazin-1-ylpropoxy, 2-homopiperazin-1-ylethoxy or 3-homopiperazin-1-ylpropoxy, provided that, any compound wherein  $R^2$  is a 2-piperidinoethoxy, 3-piperidinopropoxy, 2-morpholinoethoxy, 3-morpholinopropoxy, 2-piperazin-1-ylethoxy or  
 30 3-piperazin-1-ylpropoxy group that is unsubstituted on the piperidino, morpholino or piperazin-1-yl group or is unsubstituted on the ethoxy or propoxy group is excluded, and provided that any compound wherein  $R^2$  is a 2-[4-(1-4C)alkyl]piperazin-1-ylethoxy or 3-[4-(1-4C)alkyl]piperazin-1-ylpropoxy group that bears no further substituent on the

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4-(1-4C)alkylpiperazin-1-yl group or is unsubstituted on the ethoxy or propoxy group is also excluded,

or R<sup>2</sup> is a group selected from 3-aminopropoxy, 3-methylaminopropoxy, 3-ethylaminopropoxy, 3-isopropylaminopropoxy, 3-dimethylaminopropoxy, 3-diethylaminopropoxy or 3-(N-isopropyl-N-methylamino)propoxy, provided that the central CH<sub>2</sub> group within the propoxy group bears a hydroxy or acetoxy substituent,

and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a R<sup>2</sup> substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, cyano, amino, methoxy, methylsulphonyl, methylamino and dimethylamino,

and wherein any pyridyl or heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, cyano, hydroxy, amino, methyl, ethyl and methoxy,

and wherein any heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 oxo substituents; and

R<sup>3</sup> is chloro or bromo;  
or a pharmaceutically-acceptable acid-addition salt thereof.

3. A quinazoline derivative of the Formula I according to claim 1 wherein :

R<sup>1</sup> is methoxy and R<sup>2</sup> is 2-imidazol-1-ylethoxy, 3-(1,2,3-triazol-1-yl)propoxy, 4-pyridylmethoxy, 2-pyrid-4-yloxyethoxy, 3-pyrid-2-yloxypropoxy, 3-pyrid-4-yloxypropoxy, 3-azetidin-1-yl-2-hydroxypropoxy, 2-pyrrolidin-1-ylethoxy, 3-pyrrolidin-1-ylpropoxy, 2-hydroxy-3-pyrrolidin-1-ylpropoxy, pyrrolidin-3-yloxy, pyrrolidin-2-ylmethoxy, 2-pyrrolidin-2-ylethoxy, 3-pyrrolidin-2-ylpropoxy, 2-hydroxy-3-morpholinopropoxy, 2-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)ethoxy, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-hydroxy-3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-hydroxy-3-piperidinopropoxy, piperidin-3-yloxy, piperidin-4-yloxy, piperidin-3-ylmethoxy, 2-piperidin-3-ylethoxy, piperidin-4-ylmethoxy, 2-piperidin-4-ylethoxy, 3-homopiperidin-1-ylpropoxy, 3-homopiperidin-1-yl-2-hydroxypropoxy, 2-hydroxy-3-piperazin-1-ylpropoxy, 3-homopiperazin-1-ylpropoxy or 2-hydroxy-3-homopiperazin-1-ylpropoxy,

or R<sup>2</sup> is a group selected from 2-hydroxy-3-methylaminopropoxy, 3-ethylamino-2-hydroxypropoxy, 2-hydroxy-3-isopropylaminopropoxy, 3-dimethylamino-2-hydroxypropoxy, 3-diethylamino-2-hydroxypropoxy, 3-(N-ethyl-

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N-isopropylamino)-2-hydroxypropoxy, 3-(N-ethyl-N-methylamino)-2-hydroxypropoxy or 3-(N-isopropyl-N-methylamino)-2-hydroxypropoxy,

and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a R<sup>2</sup> substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, cyano, amino, vinyl, ethynyl, methoxy, methylsulphonyl, methylamino and dimethylamino, and wherein any heteroaryl or heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, cyano, hydroxy, amino, methyl, ethyl, cyclopropyl, allyl, methoxy and acetyl, and wherein any heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 oxo substituents; and

R<sup>3</sup> is chloro or bromo;

or a pharmaceutically-acceptable acid-addition salt thereof.

4. A quinazoline derivative of the Formula I according to claim 1 wherein :

R<sup>1</sup> is methoxy and R<sup>2</sup> is 4-pyridylmethoxy, 2-pyrid-4-yloxyethoxy, 3-pyrid-2-yloxypropoxy, 3-pyrid-4-yloxypropoxy, 3-azetidin-1-yl-2-hydroxypropoxy, 2-pyrrolidin-1-ylethoxy, 3-pyrrolidin-1-ylpropoxy, 2-hydroxy-3-pyrrolidin-1-ylpropoxy, 2-hydroxy-3-morpholinopropoxy, 3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-hydroxy-3-(1,1-dioxotetrahydro-4H-1,4-thiazin-4-yl)propoxy, 2-hydroxy-3-piperidinopropoxy, piperidin-4-ylmethoxy, 2-piperidin-4-ylethoxy, 3-homopiperidin-1-yl-2-hydroxypropoxy or 2-hydroxy-3-piperazin-1-ylpropoxy,

or R<sup>2</sup> is a group selected from 3-dimethylamino-2-hydroxypropoxy, 3-diethylamino-2-hydroxypropoxy, 3-(N-ethyl-N-isopropylamino)-2-hydroxypropoxy, 3-(N-ethyl-N-methylamino)-2-hydroxypropoxy or 3-(N-isopropyl-N-methylamino)-2-hydroxypropoxy,

and wherein any CH<sub>2</sub> or CH<sub>3</sub> group within a R<sup>2</sup> substituent optionally bears on each said CH<sub>2</sub> or CH<sub>3</sub> group a substituent selected from hydroxy, cyano, amino, vinyl, ethynyl, methoxy, methylsulphonyl, methylamino and dimethylamino,

and wherein any pyridyl or heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 substituents, which may be the same or different, selected from fluoro, chloro, trifluoromethyl, cyano, hydroxy, amino, methyl, ethyl, cyclopropyl, allyl, methoxy and acetyl, and wherein any heterocyclyl group within a substituent on R<sup>2</sup> optionally bears 1 or 2 oxo substituents; and

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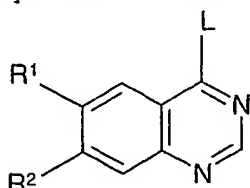
$R^3$  is chloro or bromo;  
or a pharmaceutically-acceptable acid-addition salt thereof.

5. A quinazoline derivative of the Formula I according to claim 1 selected from :

- 5 4-(2-chloro-5-methoxyanilino)-6-methoxy-7-(N-methylpiperidin-4-ylmethoxy)quinazoline,  
4-(2-chloro-5-methoxyanilino)-6-methoxy-7-piperidin-4-ylmethoxyquinazoline and  
4-(2-bromo-5-methoxyanilino)-6-methoxy-7-[2-(N-methylpiperidin-4-yl)ethoxy]quinazoline;  
or a pharmaceutically-acceptable acid-addition salt thereof.

- 10 6. A process for the preparation of a quinazoline derivative of the Formula I, or a  
pharmaceutically-acceptable salt thereof, according to claim 1 which comprises :-

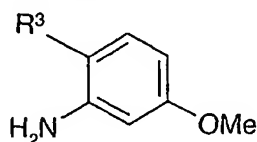
(a) the reaction of a quinazoline of the Formula II



II

wherein L is a displaceable group and  $R^1$  and  $R^2$  have any of the meanings defined in claim 1

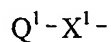
- 15 except that any functional group is protected if necessary, with an aniline of the Formula III



III

wherein  $R^3$  has any of the meanings defined in claim 1 except that any functional group is  
protected if necessary, whereafter any protecting group that is present is removed by  
conventional means;

- 20 (b) for the production of those compounds of the Formula I wherein  $R^2$  is a group of the  
formula :

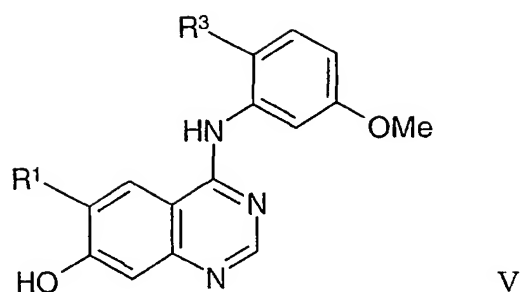


wherein  $X^1$  is an oxygen atom, the coupling of an alcohol of the Formula



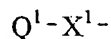
- 25 wherein  $Q^1$  has any of the meanings defined in claim 1 except that any functional group is  
protected if necessary, with a quinazoline of the Formula V

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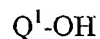


wherein  $R^1$  and  $R^3$  have any of the meanings defined in claim 1 except that any functional group is protected if necessary, whereafter any protecting group that is present is removed by conventional means;

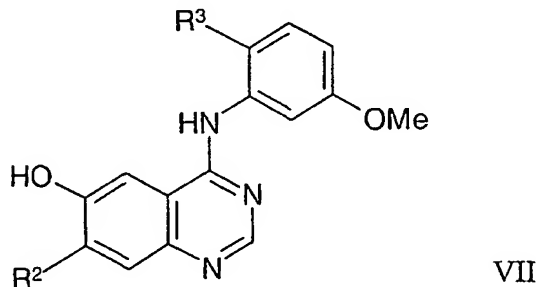
- 5 (c) for the production of those compounds of the Formula I wherein  $R^1$  is a group of the formula :



wherein  $X^1$  is an oxygen atom, the coupling of an alcohol of the Formula



- 10 wherein  $Q^1$  has any of the meanings defined in claim 1 except that any functional group is protected if necessary, with a quinazoline of the Formula VII



wherein  $R^2$  and  $R^3$  have any of the meanings defined in claim 1 except that any functional group is protected if necessary, whereafter any protecting group that is present is removed by

- 15 conventional means;

(d) for the production of those compounds of the Formula I wherein  $R^1$  or  $R^2$  contains an amino-hydroxy-disubstituted (1-6C)alkoxy group, the reaction of a compound of the Formula I wherein  $R^1$  or  $R^2$  contains an epoxy-substituted (1-6C)alkoxy group with a heterocyclyl compound or an appropriate amine;

- 20 (e) for the production of those compounds of the Formula I wherein  $R^1$  or  $R^2$  contains an amino-acyloxy-disubstituted (1-6C)alkoxy group, the acylation of a compound of the Formula I wherein  $R^1$  or  $R^2$  contains an amino-hydroxy-disubstituted (1-6C)alkoxy group; or

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(f) for the production of those compounds of the Formula I wherein an R<sup>1</sup> or R<sup>2</sup> group contains a hydroxy group, the cleavage of the corresponding compound of the Formula I wherein the R<sup>1</sup> or R<sup>2</sup> group contains a protected hydroxy group;

and when a pharmaceutically-acceptable salt of a quinazoline derivative of the  
5 Formula I is required, it may be obtained using a conventional procedure.

7. A pharmaceutical composition which comprises a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, according to claim 1 in association with a pharmaceutically-acceptable diluent or carrier.

10

8. A quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, according to claim 1 for use in a method of treatment of the human or animal body by therapy.

15 9. The use of a quinazoline derivative of the Formula I, or a pharmaceutically-acceptable salt thereof, according to claim 1 in the manufacture of a medicament for use as an anti-invasive agent in the containment and/or treatment of solid tumour disease.



## INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 02/02117

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D239/94 C07D401/12 C07D403/12 C07D417/12 A61K31/505  
A61P35/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|------------|--|-----------------------|
| Y          | EP 0 566 226 A (ZENECA)<br>20 October 1993 (1993-10-20)<br>claims<br>---           | 1,6-9                 |
| Y          | WO 97 32856 A (ZENECA)<br>12 September 1997 (1997-09-12)<br>claims<br>---          | 1,6-9                 |
| Y          | WO 98 13354 A (ZENECA)<br>2 April 1998 (1998-04-02)<br>claims<br>---               | 1,6-9                 |
|            | ---<br>-/--  |                       |

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

## \* Special categories of cited documents:

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\*B\* document member of the same patent family

Date of the actual completion of the international search

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PCT/GB 02/02117

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages   | Relevant to claim No. |
|------------|--|-----------------------|
| P, Y       | DIANE BOSCHELLI ET AL.: "OPTIMISATION OF<br>4-PHENYLAMINO-3-QUINOLINECARBONITRILE AS<br>POTENT INHIBITORS OF SRC KINASE ACTIVITY."<br>JOURNAL OF MEDICINAL CHEMISTRY.,<br>vol. 44, no. 23, 2001, pages 3965-77,<br>XP002211143<br>AMERICAN CHEMICAL SOCIETY. WASHINGTON., US<br>ISSN: 0022-2623<br>page 3965 -page 3972; example 3B<br>----- | 1, 6-9                |

## INTERNATIONAL SEARCH REPORT

Information on patent family members

national Application No

PCT/GB 02/02117

| Patent document<br>cited in search report |   | Publication<br>date | Patent family<br>member(s) | Publication<br>date |
|---|---|---------------------|----------------------------|---------------------|
| EP 566226                                 | A | 20-10-1993          | AT 130000 T                | 15-11-1995          |
|   |   |                     | AU 661533 B2               | 27-07-1995          |
|   |   |                     | AU 3101093 A               | 22-07-1993          |
|   |   |                     | CA 2086968 A1              | 21-07-1993          |
|   |   |                     | CZ 9300043 A3              | 13-10-1993          |
|   |   |                     | DE 69300754 D1             | 14-12-1995          |
|   |   |                     | DE 69300754 T2             | 28-03-1996          |
|   |   |                     | DK 566226 T3               | 18-03-1996          |
|   |   |                     | EP 0566226 A1              | 20-10-1993          |
|   |   |                     | ES 2078798 T3              | 16-12-1995          |
|   |   |                     | FI 930208 A                | 21-07-1993          |
|   |   |                     | GR 3018143 T3              | 29-02-1996          |
|   |   |                     | HK 36497 A                 | 04-04-1997          |
|   |   |                     | HU 63153 A2                | 28-07-1993          |
|   |   |                     | HU 9500185 A3              | 28-07-1995          |
|   |   |                     | IL 104479 A                | 22-12-1999          |
|   |   |                     | KR 229294 B1               | 01-11-1999          |
|   |   |                     | MX 9300277 A1              | 30-06-1994          |
|   |   |                     | NO 930178 A                | 21-07-1993          |
|   |   |                     | NZ 245662 A                | 26-09-1995          |
|   |   |                     | RU 2127263 C1              | 10-03-1999          |
|   |   |                     | SK 1693 A3                 | 09-09-1993          |
|   |   |                     | US 5457105 A               | 10-10-1995          |
|   |   |                     | US 5616582 A               | 01-04-1997          |
|   |   |                     | ZA 9300015 A               | 20-07-1993          |
|   |   |                     | JP 2994165 B2              | 27-12-1999          |
|   |   |                     | JP 6073025 A               | 15-03-1994          |
| WO 9732856                                | A | 12-09-1997          | AT 211134 T                | 15-01-2002          |
|   |   |                     | AU 719327 B2               | 04-05-2000          |
|   |   |                     | AU 1866497 A               | 22-09-1997          |
|   |   |                     | CA 2244897 A1              | 12-09-1997          |
|   |   |                     | CN 1212684 A               | 31-03-1999          |
|   |   |                     | DE 69709319 D1             | 31-01-2002          |
|   |   |                     | DE 69709319 T2             | 14-08-2002          |
|   |   |                     | DK 885198 T3               | 25-03-2002          |
|   |   |                     | EP 0885198 A1              | 23-12-1998          |
|   |   |                     | ES 2169355 T3              | 01-07-2002          |
|   |   |                     | WO 9732856 A1              | 12-09-1997          |
|   |   |                     | JP 2000517291 T            | 26-12-2000          |
|   |   |                     | NO 984085 A                | 04-09-1998          |
|   |   |                     | NZ 331191 A                | 27-03-2000          |
|   |   |                     | US 6291455 B1              | 18-09-2001          |
|   |   |                     | ZA 9701747 A               | 27-08-1998          |
| WO 9813354                                | A | 02-04-1998          | AU 729968 B2               | 15-02-2001          |
|   |   |                     | AU 4561397 A               | 17-04-1998          |
|   |   |                     | BR 9711302 A               | 17-08-1999          |
|   |   |                     | CN 1231662 A               | 13-10-1999          |
|   |   |                     | CZ 9901039 A3              | 16-06-1999          |
|   |   |                     | EP 0929530 A1              | 21-07-1999          |
|   |   |                     | WO 9813354 A1              | 02-04-1998          |
|   |   |                     | JP 2001500891 T            | 23-01-2001          |
|   |   |                     | NO 991422 A                | 24-03-1999          |
|   |   |                     | NZ 334014 A                | 27-10-2000          |
|   |   |                     | PL 332385 A1               | 13-09-1999          |
|   |   |                     | SK 38999 A3                | 08-10-1999          |
|   |   |                     | TR 9900674 T2              | 21-07-1999          |

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No  
PCT/GB 02/02117

| Patent document<br>cited in search report | Publication<br>date | Patent family<br>member(s) | Publication<br>date |
|---|---------------------|----------------------------|---------------------|
| WO 9813354                                | A                   | US 6414148 B1              | 02-07-2002          |
|   |                     | ZA 9708553 A               | 25-03-1998          |
|   |                     | HU 9902850 A2              | 28-04-2000          |
| <hr/>                                     |                     |                            |                     |

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No  
PCT/GB 02/02117

| Patent document<br>cited in search report | Publication<br>date | Patent family<br>member(s) | Publication<br>date |
|---|---------------------|----------------------------|---------------------|
| WO 9813354                                | A                   | US 6414148 B1              | 02-07-2002          |
|   |                     | ZA 9708553 A               | 25-03-1998          |
|   |                     | HU 9902850 A2              | 28-04-2000          |
| -----                                     |                     |                            |                     |